

Chronic Kidney Disease – Mineral and Bone Disorder

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CONTINUING MEDICAL EDUCATION
DEPARTMENT OF MEDICINE



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Academic Focus

Effects of Acid on Bone
Genetic Hypercalciuria & Kidney Stone
Formation

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Disclosure Information

Employee of:	University of Rochester, Rochester, NY
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Consultant:	None
Stockholder:	Amgen

66 yo obese Black male with DM and hypertension.

BMI = 31

DM – 20 years and moderately well controlled

Hypertension – 35 years – typical BP 142/86

On:

ARB (low dose secondary to hyperkalemia)

Loop diuretic

Insulin

MVI

Vitamin D₃

GFR = 9 ml/min

K = 4.8 meq/l

Ca = 9.2 mg/dl

P = 5.6 mg/dl

25 D = 25 ng/ml

PTH = 345 pg/ml

Over the course of the next 6 months GFR falls

GFR = 4 ml/min

K = 5.3 meq/l

Ca = 9.5 mg/dl

P = 6.1 mg/dl

25 D = 22 ng/ml

PTH = 469 pg/ml

Patient has nausea, lethargy, wt loss and is started on 3X/wk HD

What do you do with respect to:

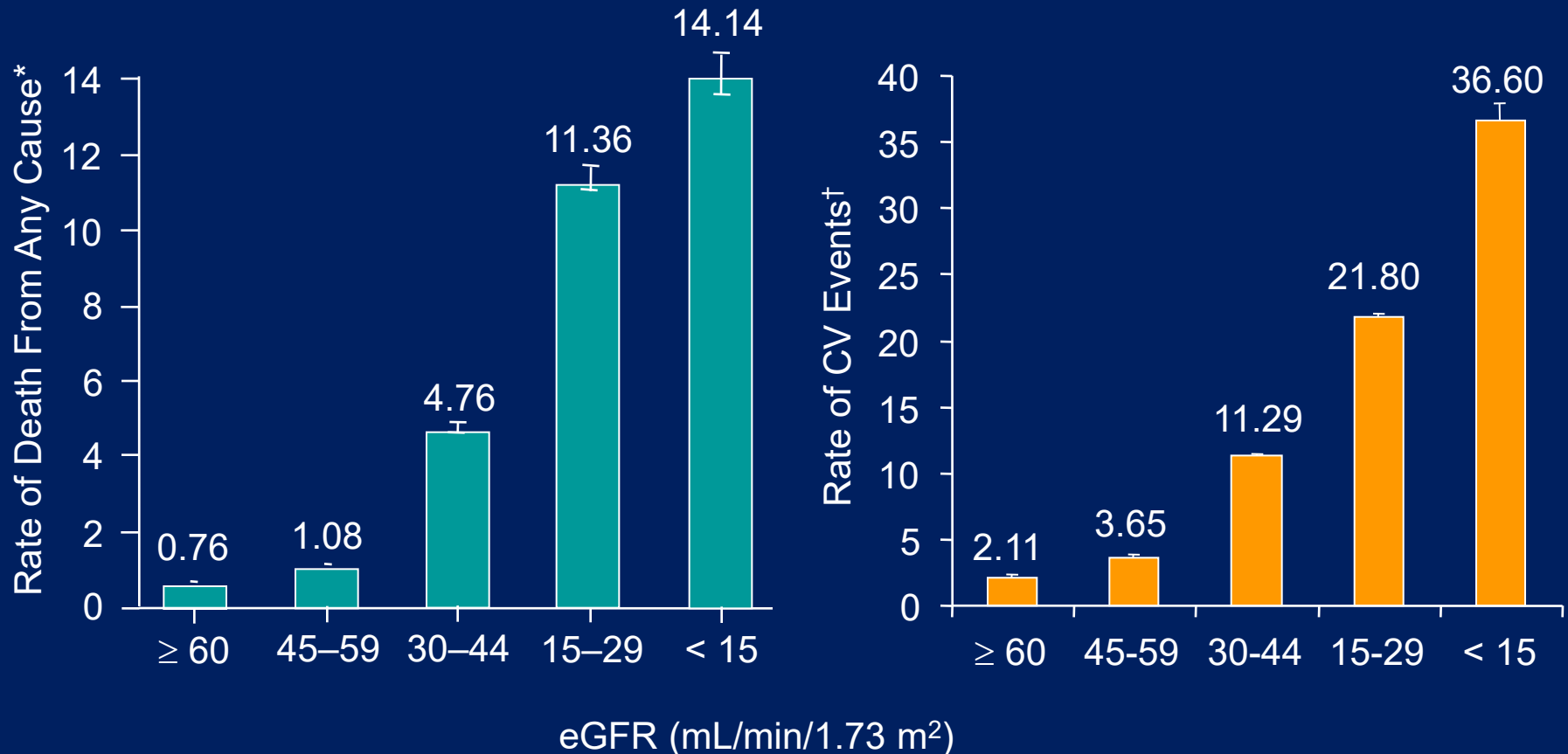
Phosphatase Binders ?

Vitamin D – dietary and/or activated ?

Cinacalcet or Etelcalcetide ?

Is he at risk for vascular calcification?
can you prevent it?

Rate of Death and Cardiovascular Events According to Estimated GFR

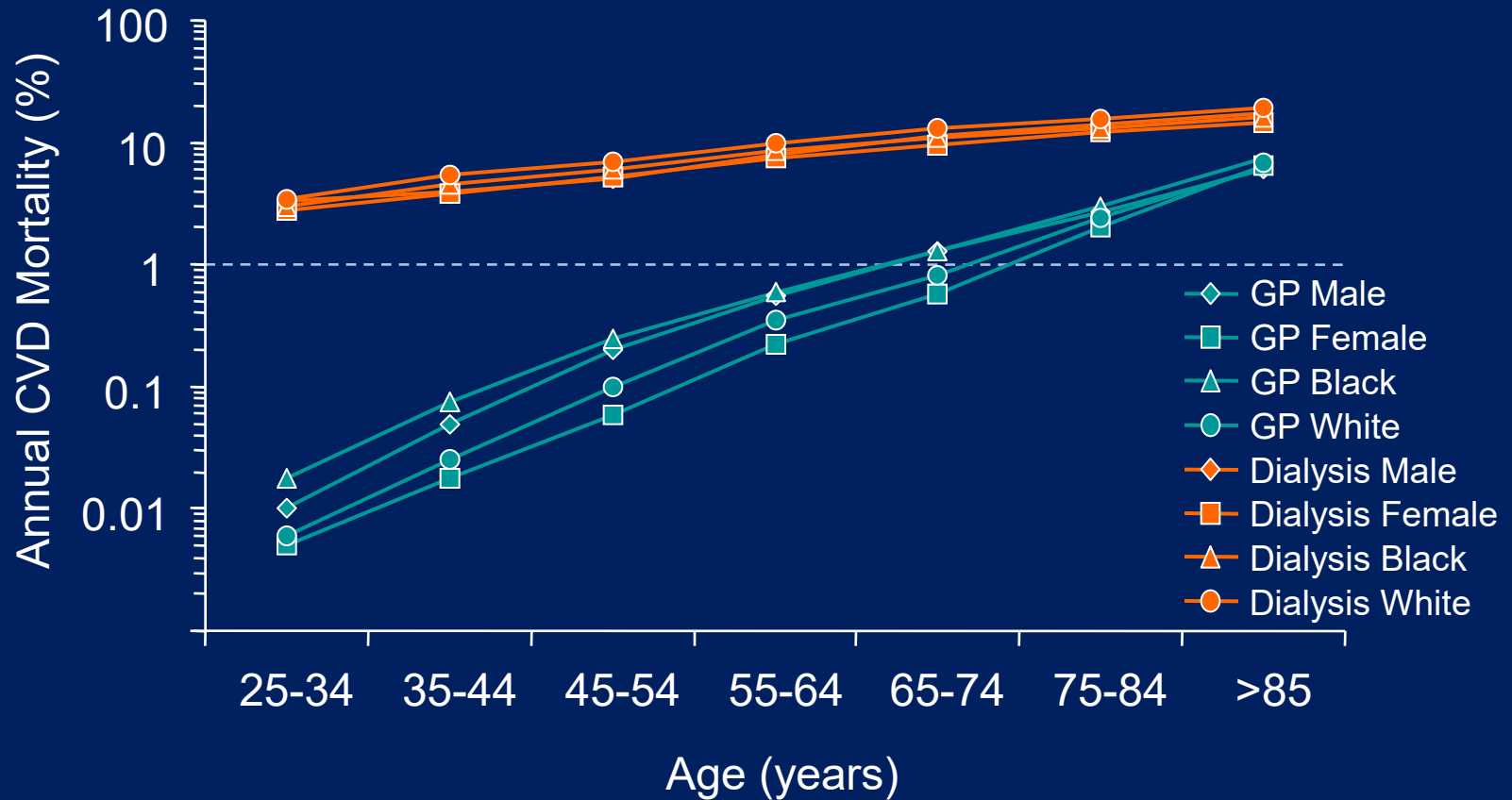


N = 1,120,295 adults.

*Age-standardized rates per 100 person-years; †CV event defined as hospitalization for coronary heart disease, heart failure, ischemic stroke, and peripheral arterial disease per 100 person-years.

Go AS, et al. *N Engl J Med*. 2004;351:1296-1305.

Dialysis Patients Have Increased Cardiovascular Disease Mortality

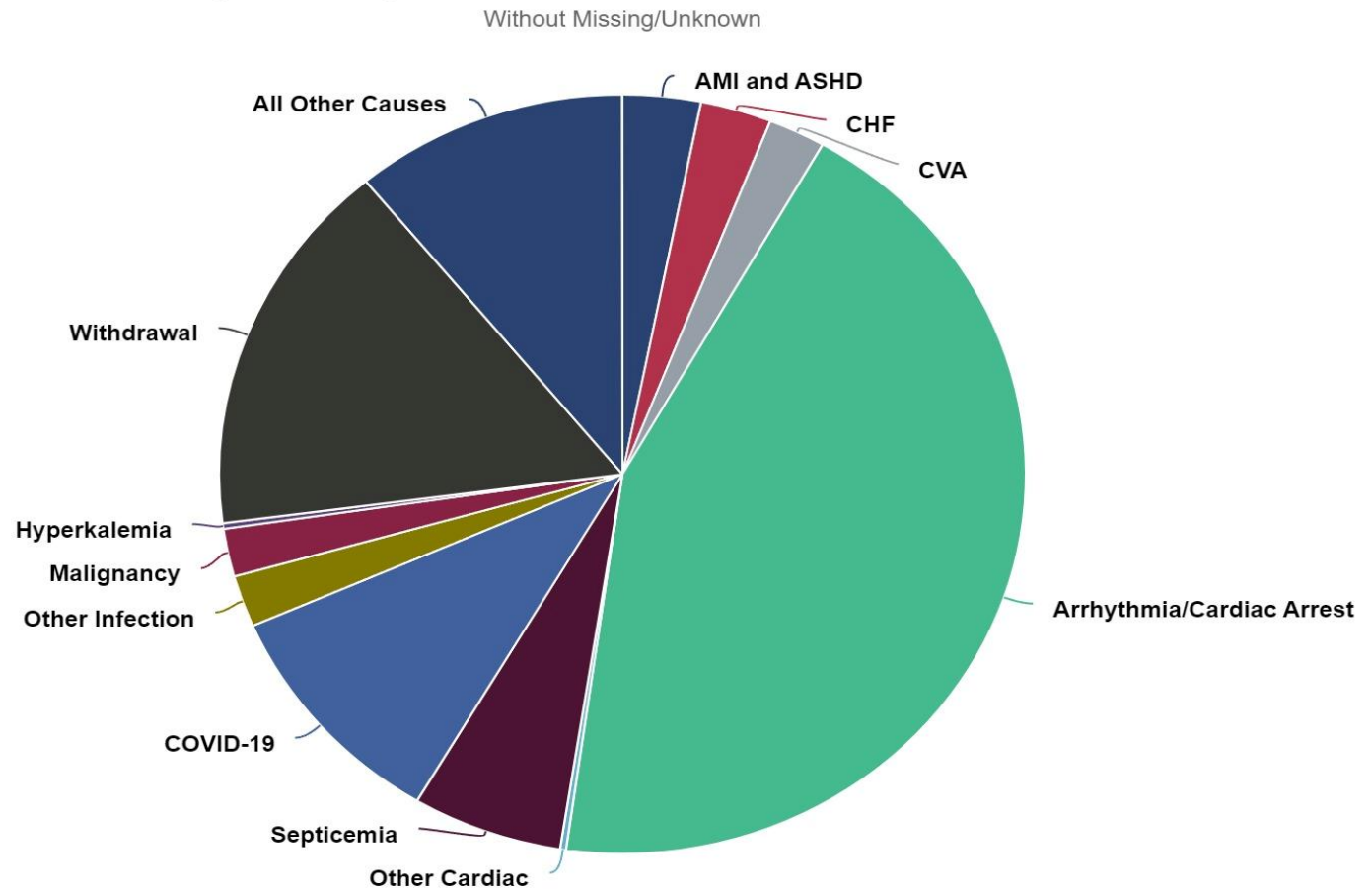


GP = general population.

Foley RN, et al. *Am J Kidney Dis*. 1998;32(suppl 3):S112-S119.

Dialysis patient mortality

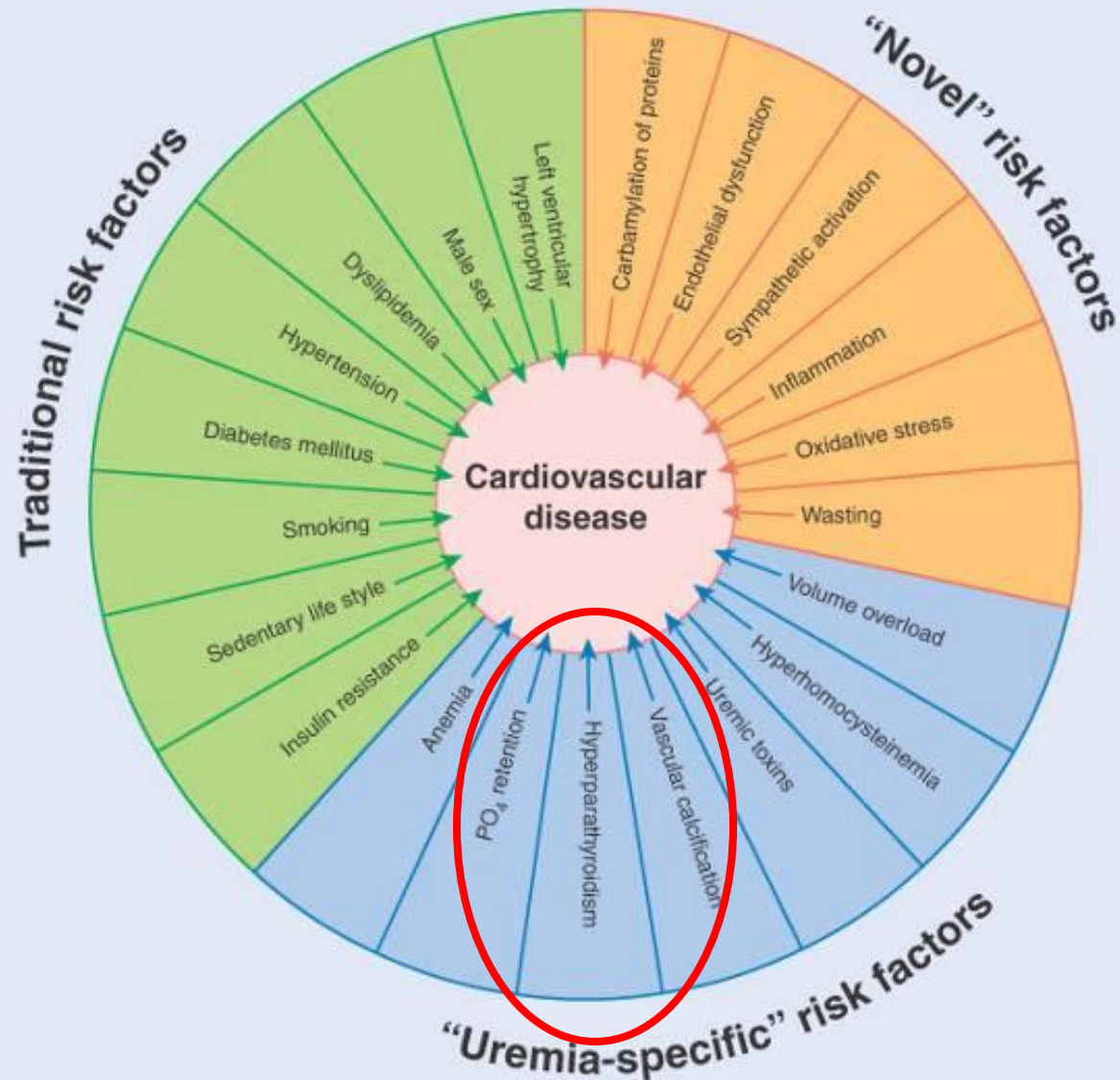
Figure 6.6a Percentages of cause-specific mortality, with and without inclusion of missing and unknown causes of death, in patients with ESRD receiving hemodialysis, who died in 2021



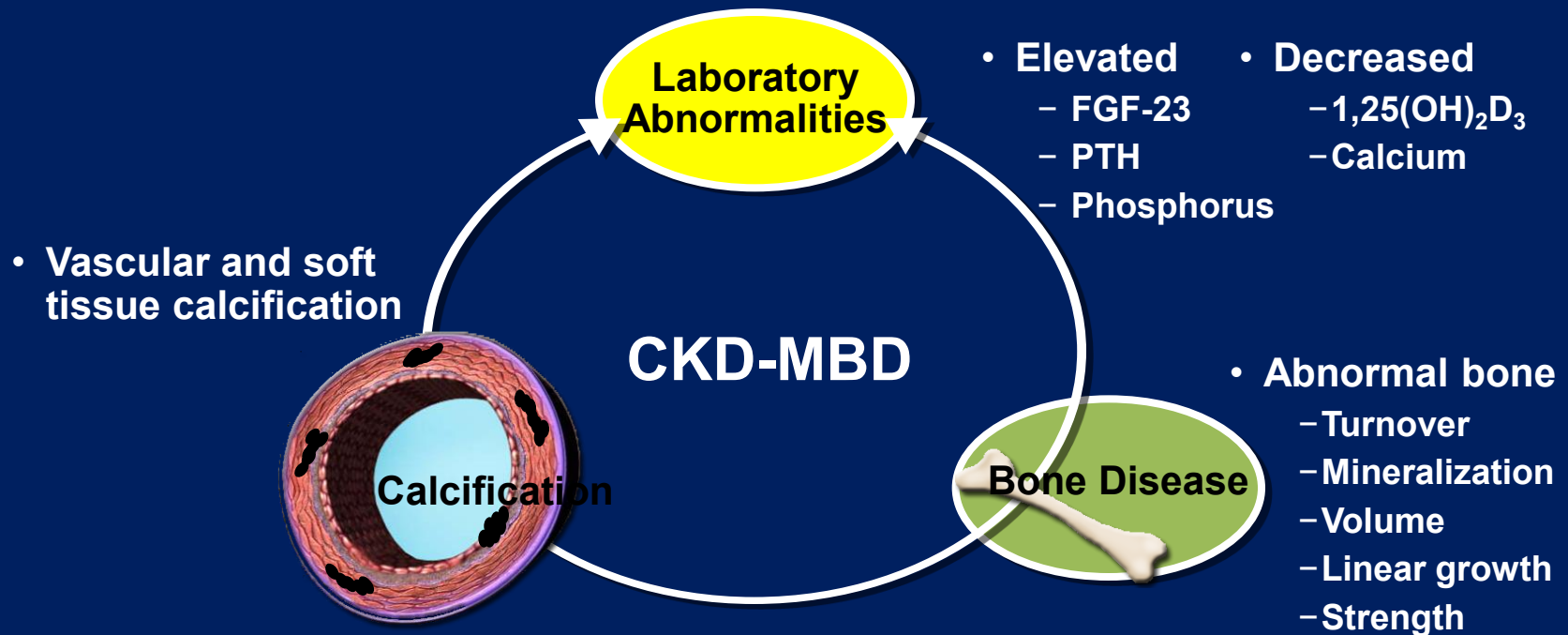
Data Source: 2023 United States Renal Data System Annual Data Report

52.5% of deaths were related to cardiovascular disease

Risk factors in chronic kidney disease



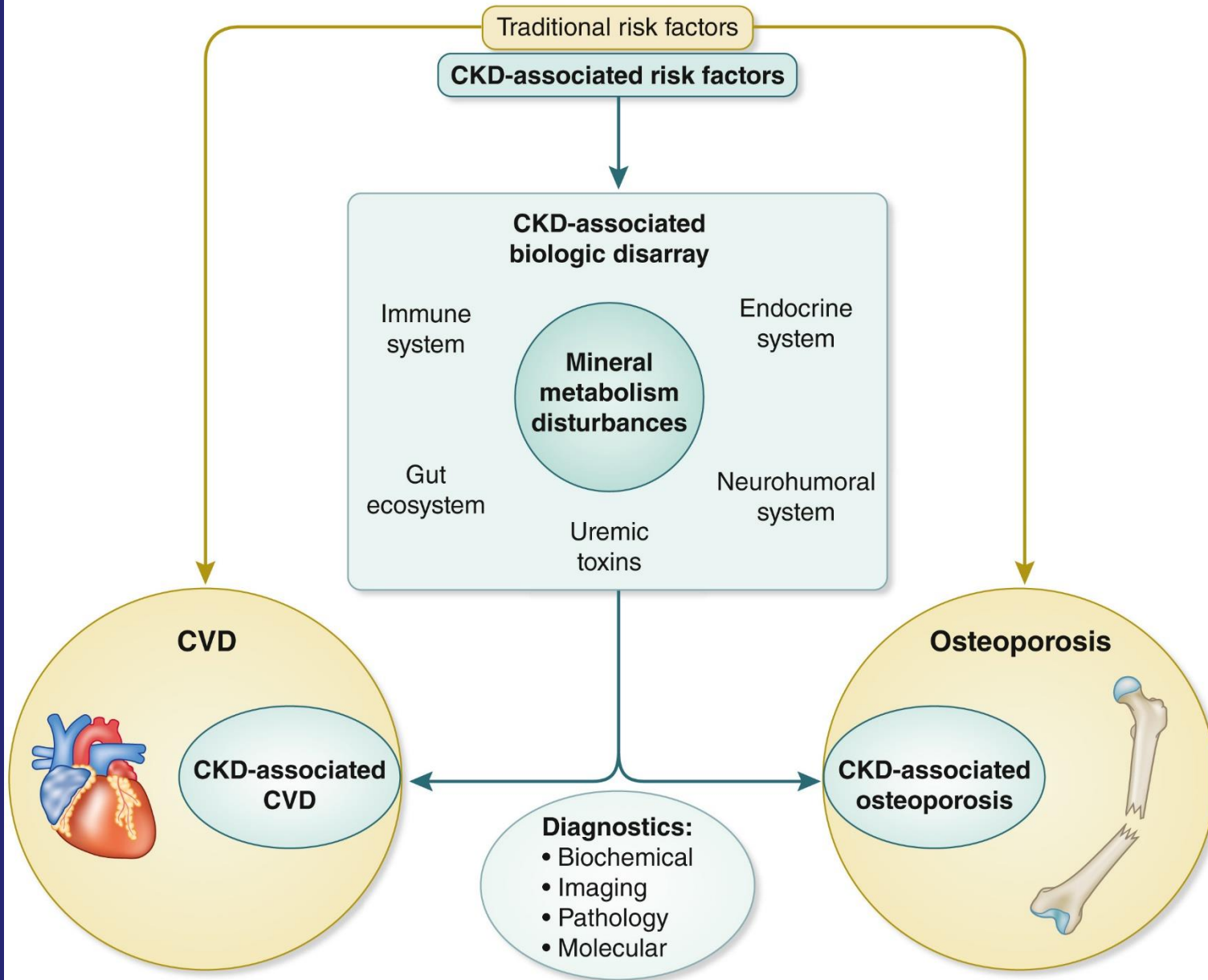
Chronic Kidney Disease-Mineral Bone Disorder



PTH = parathyroid hormone;
FGF-23 = fibroblast growth factor-23.

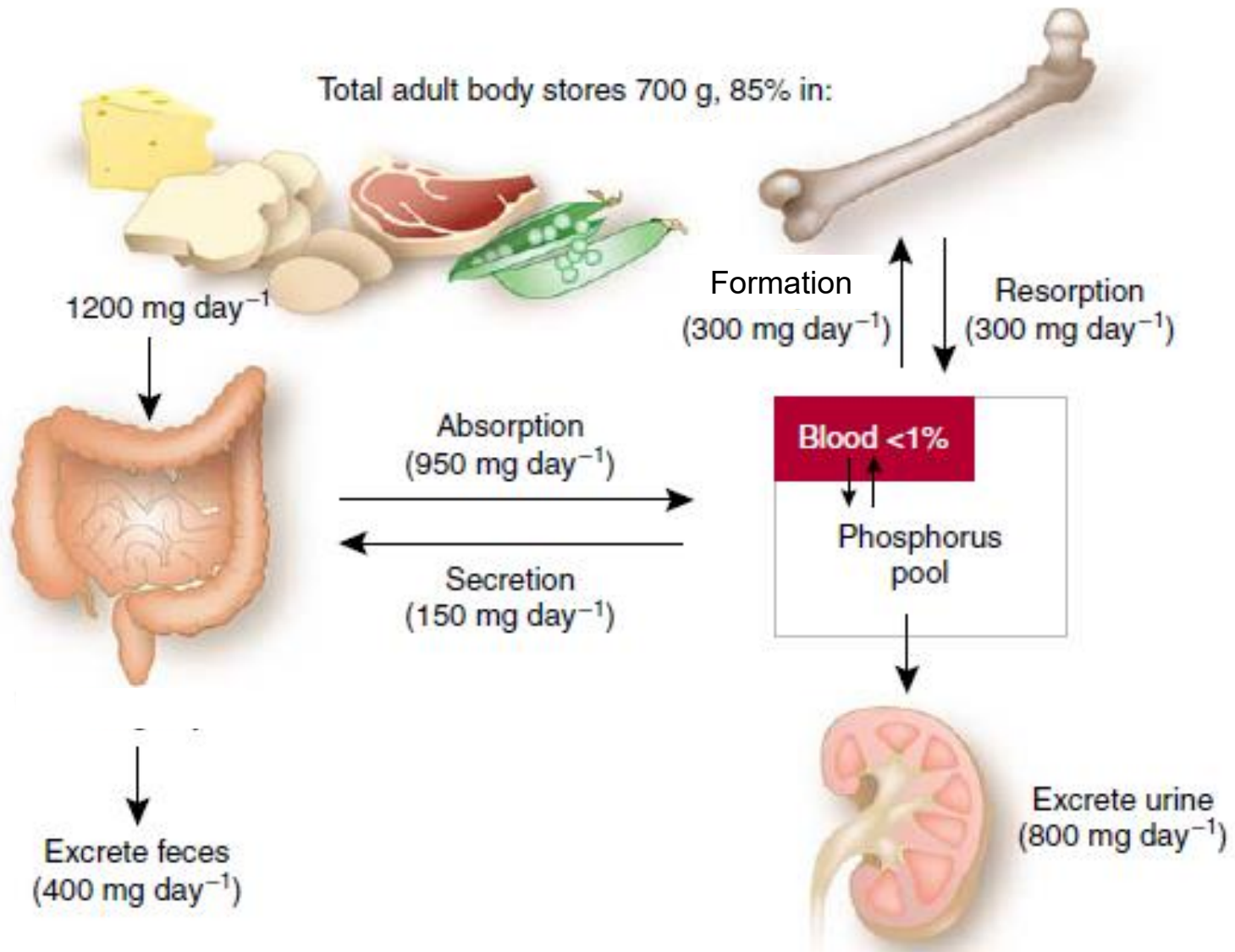
Kidney Disease: Improving Global Outcomes (KDIGO) CKD-MBD Work Group. *Kidney Int.* 2009;76

New conceptual framework moving towards personalized care in adults with CKD-MBD



Pathophysiology of CKD-MBD

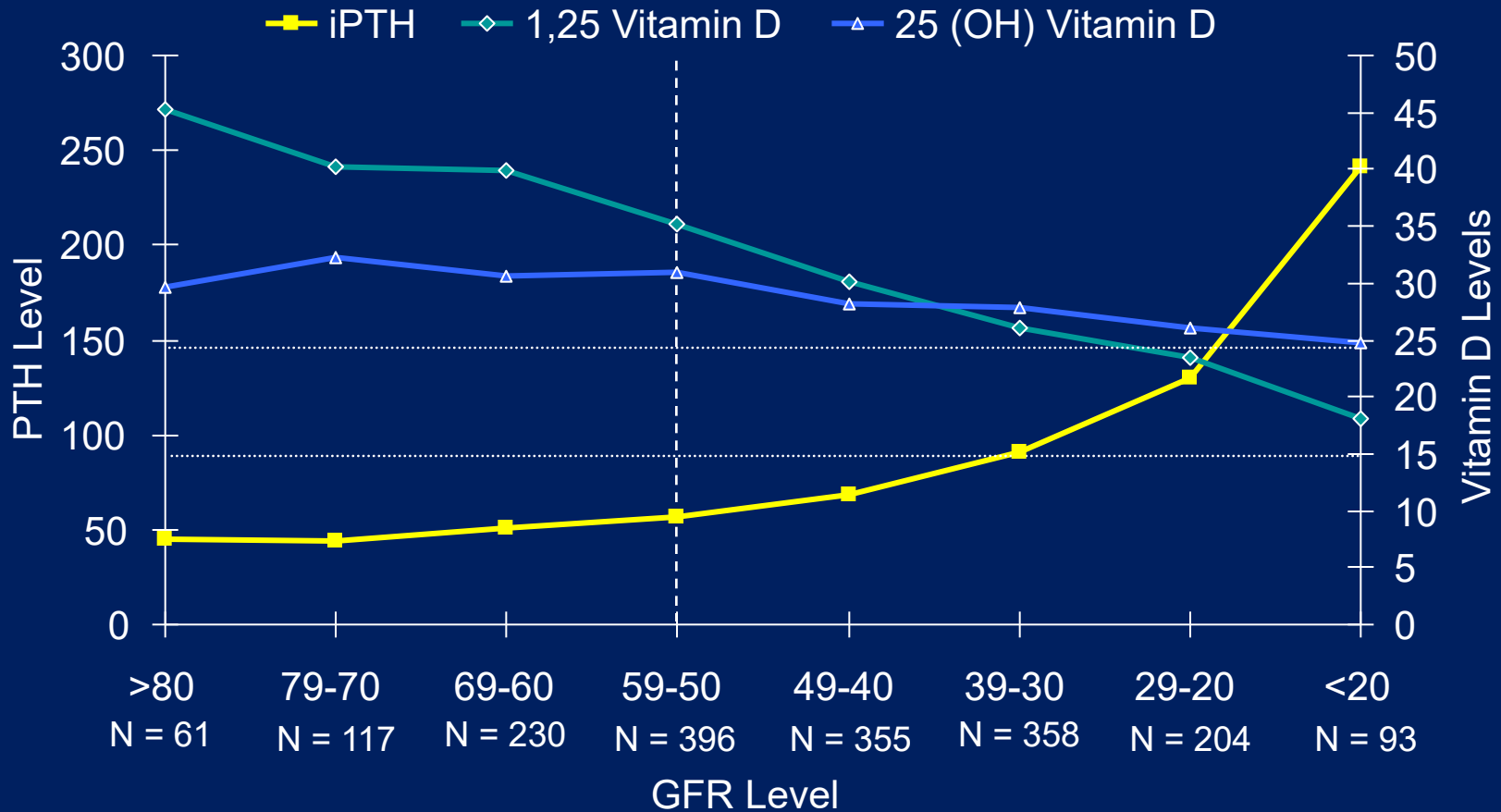
Phosphorus Balance in Health



1000 IU of oral vitamin D₃ per day will directly:

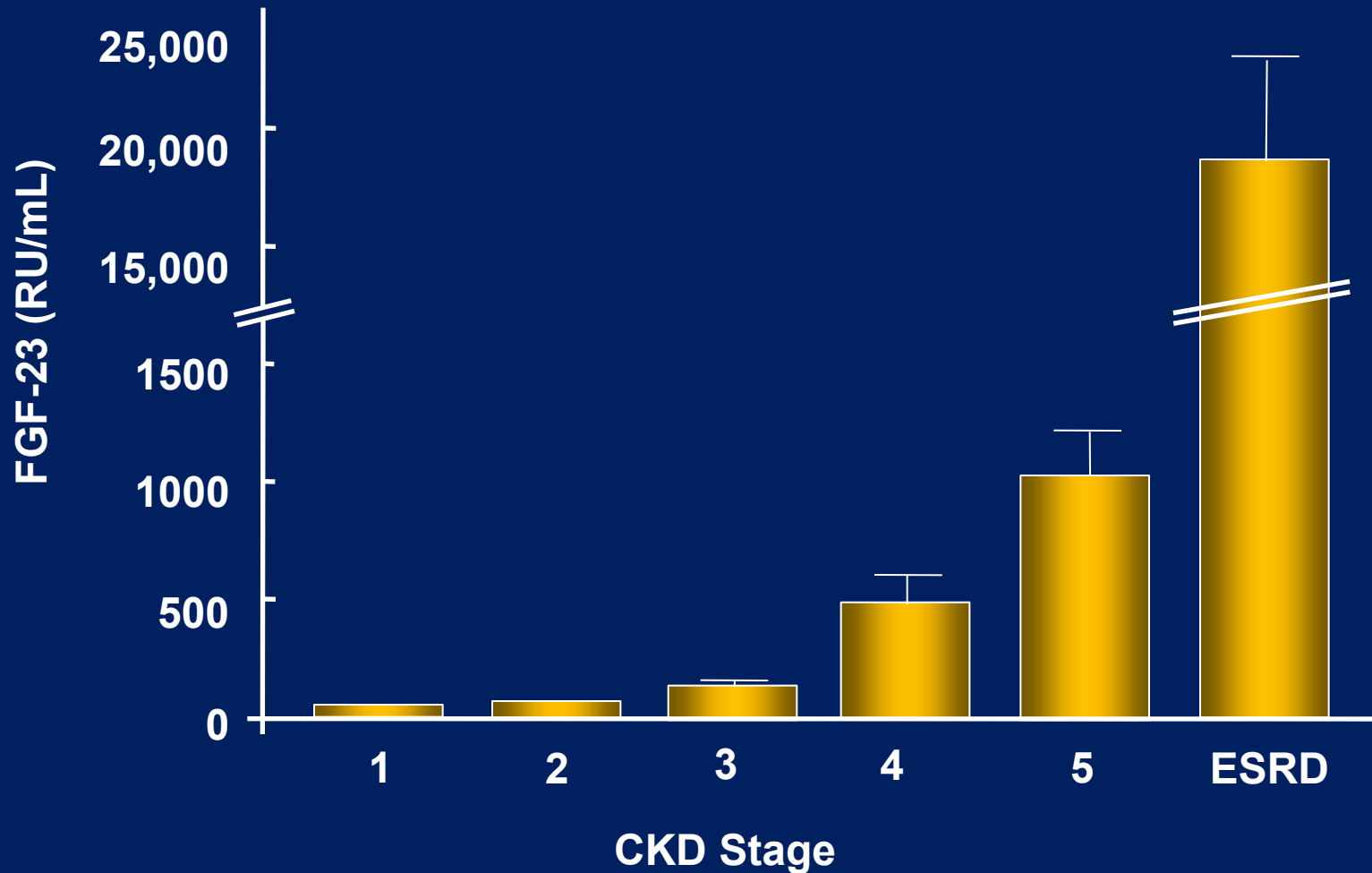
- a. increase intestinal Ca absorption, increase intestinal P absorption,
increase bone mineralization
- b. increase intestinal Ca absorption, not alter intestinal P absorption,
not alter bone mineralization
- c. not alter intestinal Ca absorption, not alter intestinal P absorption,
not alter bone mineralization
- d. not alter intestinal Ca absorption, increase intestinal P absorption,
not alter bone mineralization
- e. increase intestinal Ca absorption, not alter intestinal P absorption,
not alter bone mineralization
- f. none of the above

Progression of iPTH, Corrected Ca, and P Levels Study for the Evaluation of Early Kidney Disease (SEEK)



N = 1814

Serum FGF-23 in Various Stages of CKD

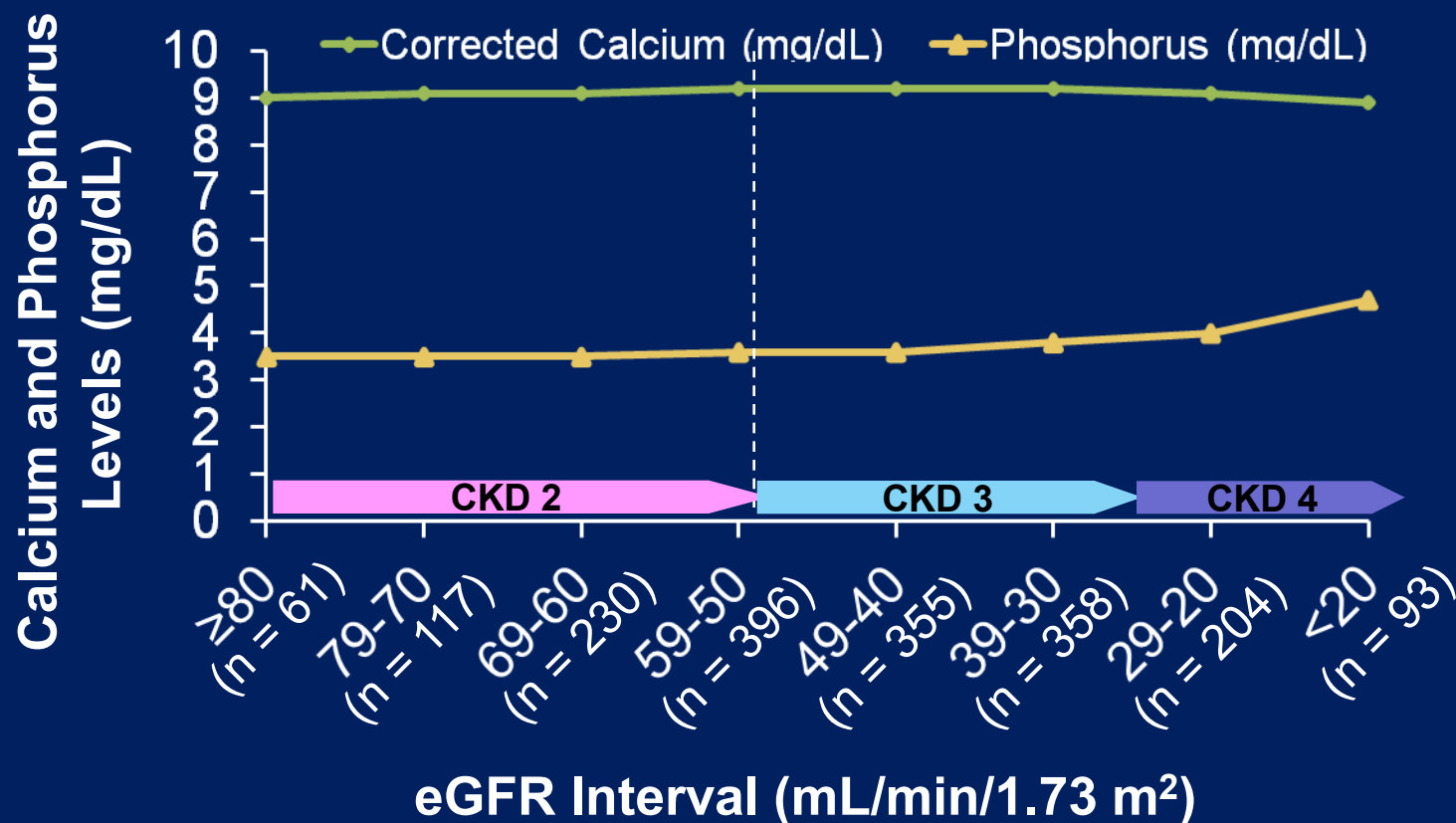


ESRD = end stage renal disease.

Pande S et al. *Nephron Physiol.* 2006;104:p23-p32.

Progression of iPTH, Corrected Ca, and P Levels Study for the Evaluation of Early Kidney Disease (SEEK)

Prospective, observational, multi-center study



N = 1814

Levin A et al. *Kidney Int.* 2007;71:31-38.

CKD-MBD

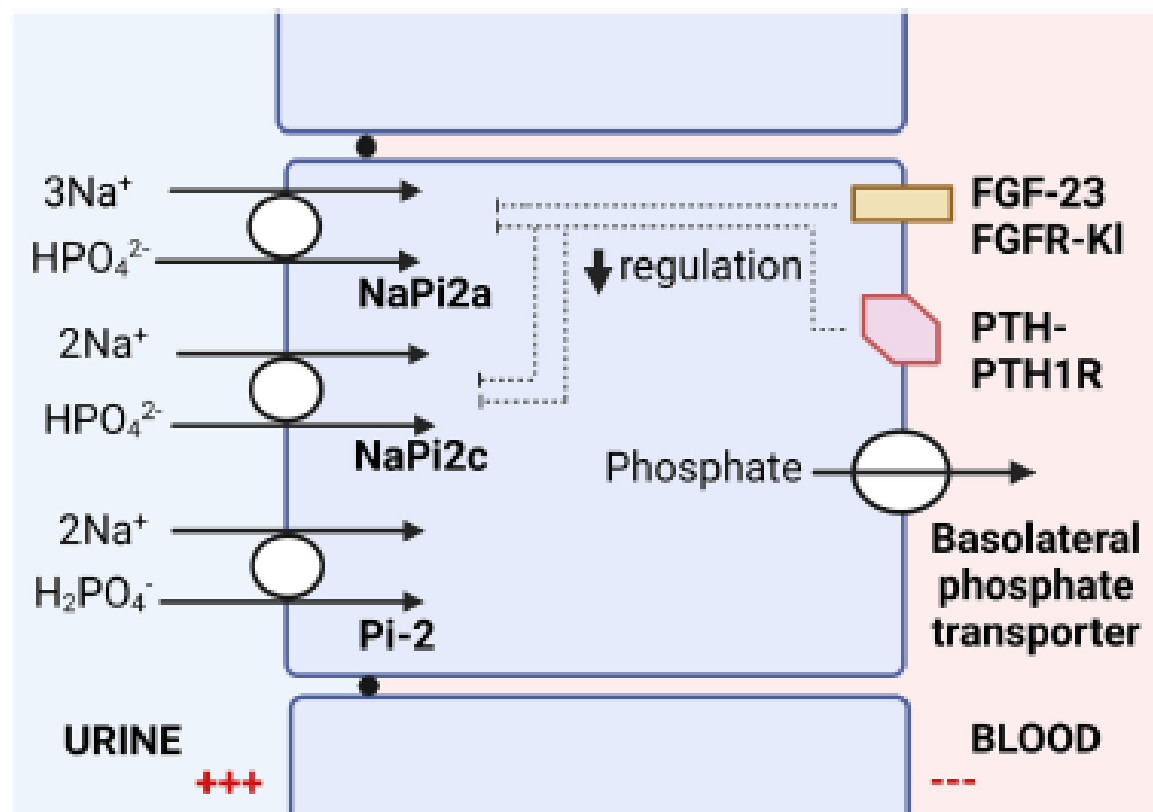


Figure 5. Phosphate transport in the proximal convoluted tubule. Phosphate is reabsorbed via transcellular transport in the proximal convoluted tubule. Three sodium-phosphate cotransporters, NaPi-2a, NaPi-2c, and PiT-2, transport phosphate from the tubular lumen into the cell. Phosphate is then transported out of the cell by an unknown basolateral phosphate transporter. Binding of FGF-23 to FGF receptor (FGFR)-Klotho (KI) complexes, and PTH to the PTH1R inhibits phosphate reabsorption by down-regulating apical expression of NaPi-2a and NaPi-2c. Created with Biorender. Abbreviations: FGF-23, fibroblast growth factor 23; PTH, parathyroid hormone; PTH1R, parathyroid hormone 1 receptor.

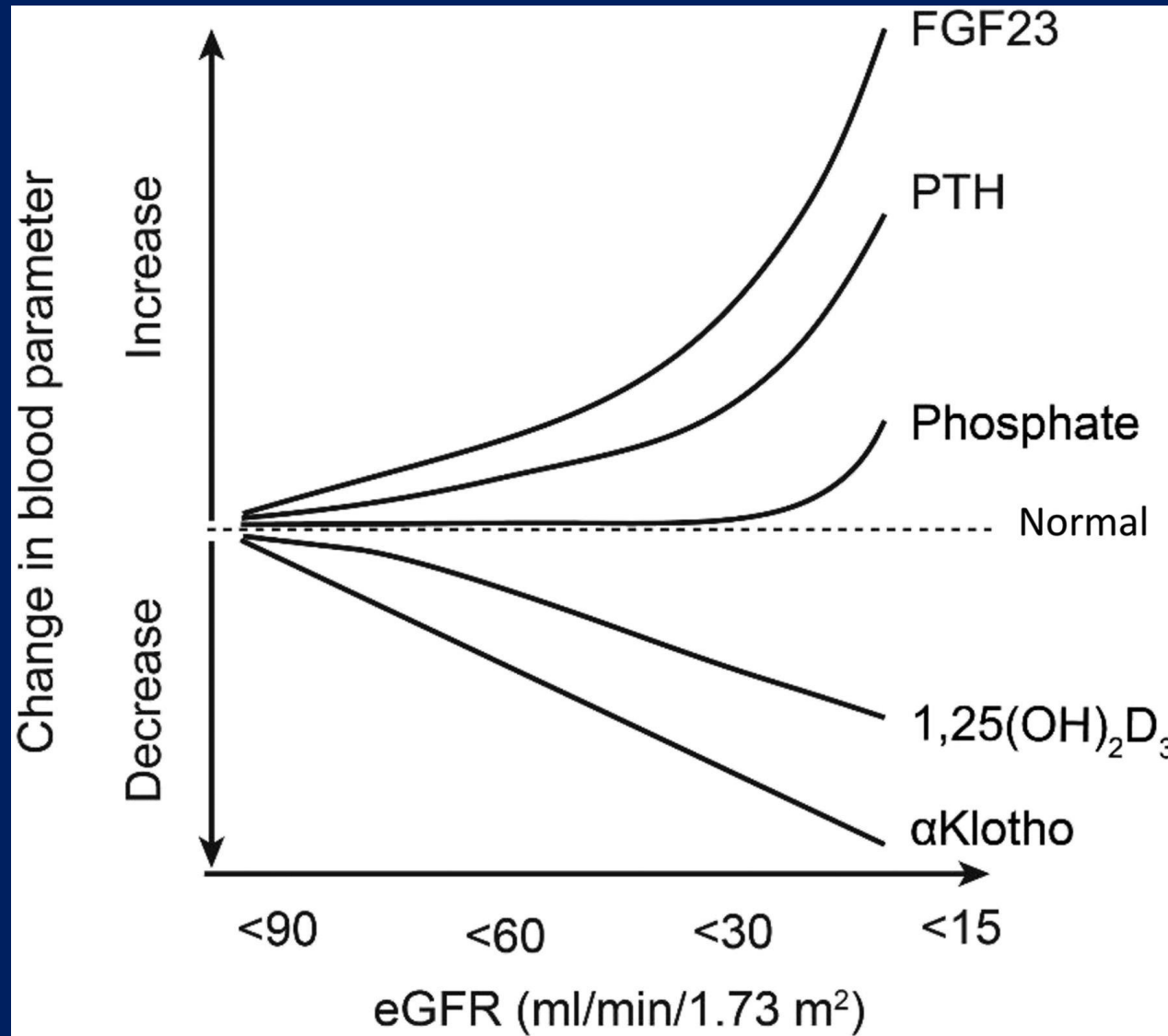
Comparison of FGF-23 and PTH

	FGF-23	PTH
Site of Production	Osteoblasts	Chief cells in parathyroid gland
Principal stimulus	Phosphorus load / hyperphosphatemia	Hypocalcemia
Principal regulator of	Phosphorus	Calcium Phosphorus
Effect on urinary phosphorus	Phosphaturic	Phosphaturic
Effect on 1,25 D production	Decreases	Increases

With respect to phosphorus, an acute infusion of FGF-23 will promptly:

- a. not alter filtered load, increase tubular reabsorption, increase urine excretion
- b. not alter filtered load, increase tubular reabsorption, stable urine excretion
- c. not alter filtered load, increase tubular reabsorption, decrease urine excretion
- d. decrease filtered load, decrease tubular reabsorption, increase urine excretion
- e. increase filtered load, decrease tubular reabsorption, increase urine excretion
- f. decrease filtered load, decrease tubular reabsorption, decrease urine excretion

Changes in circulating soluble Klotho according to eGFR decline and its relationship to other mineral metabolism parameters. 1,25(OH)₂D₃, 1,25-dihydroxyvitamin D.



CKD-MBD

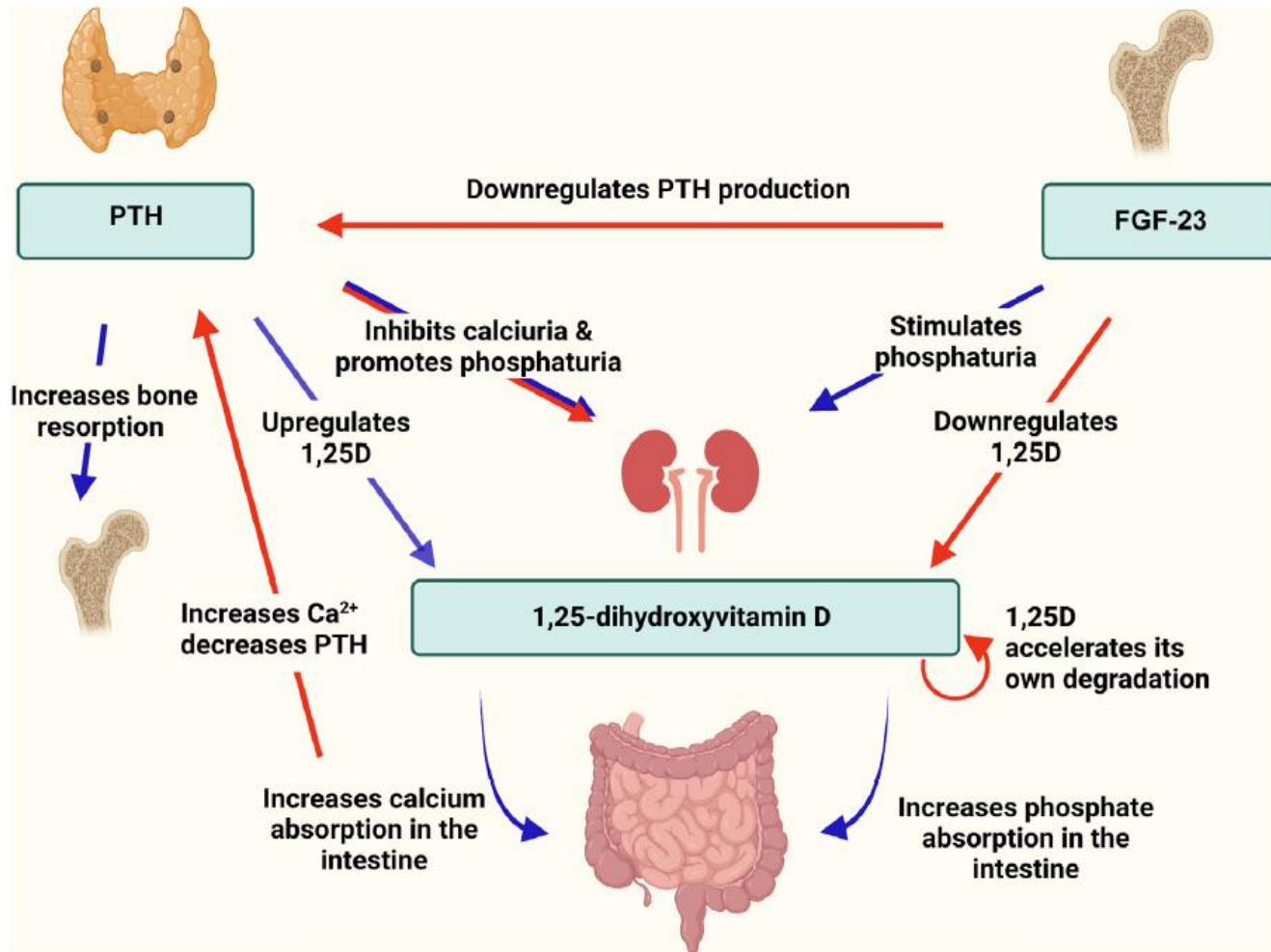
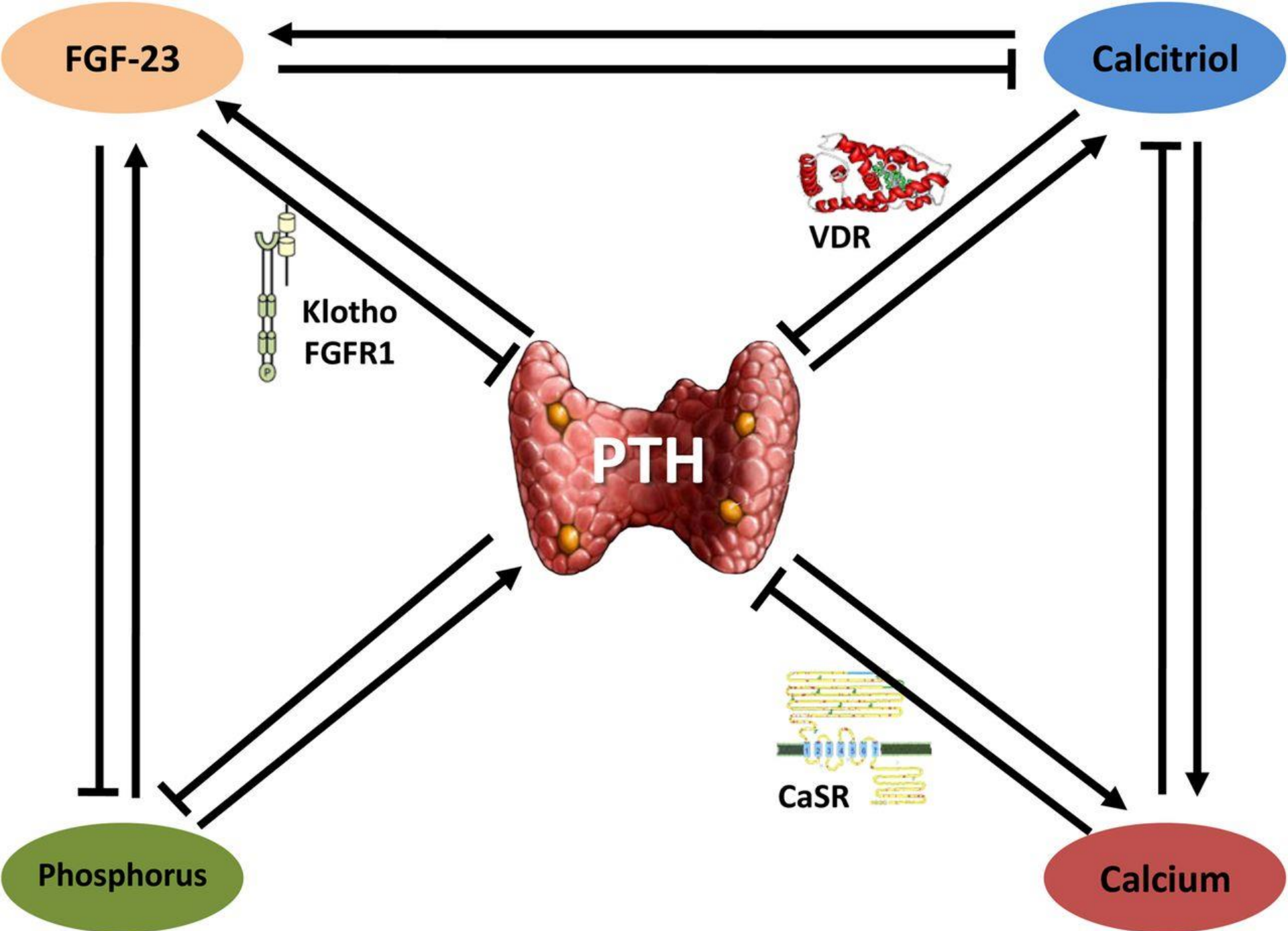
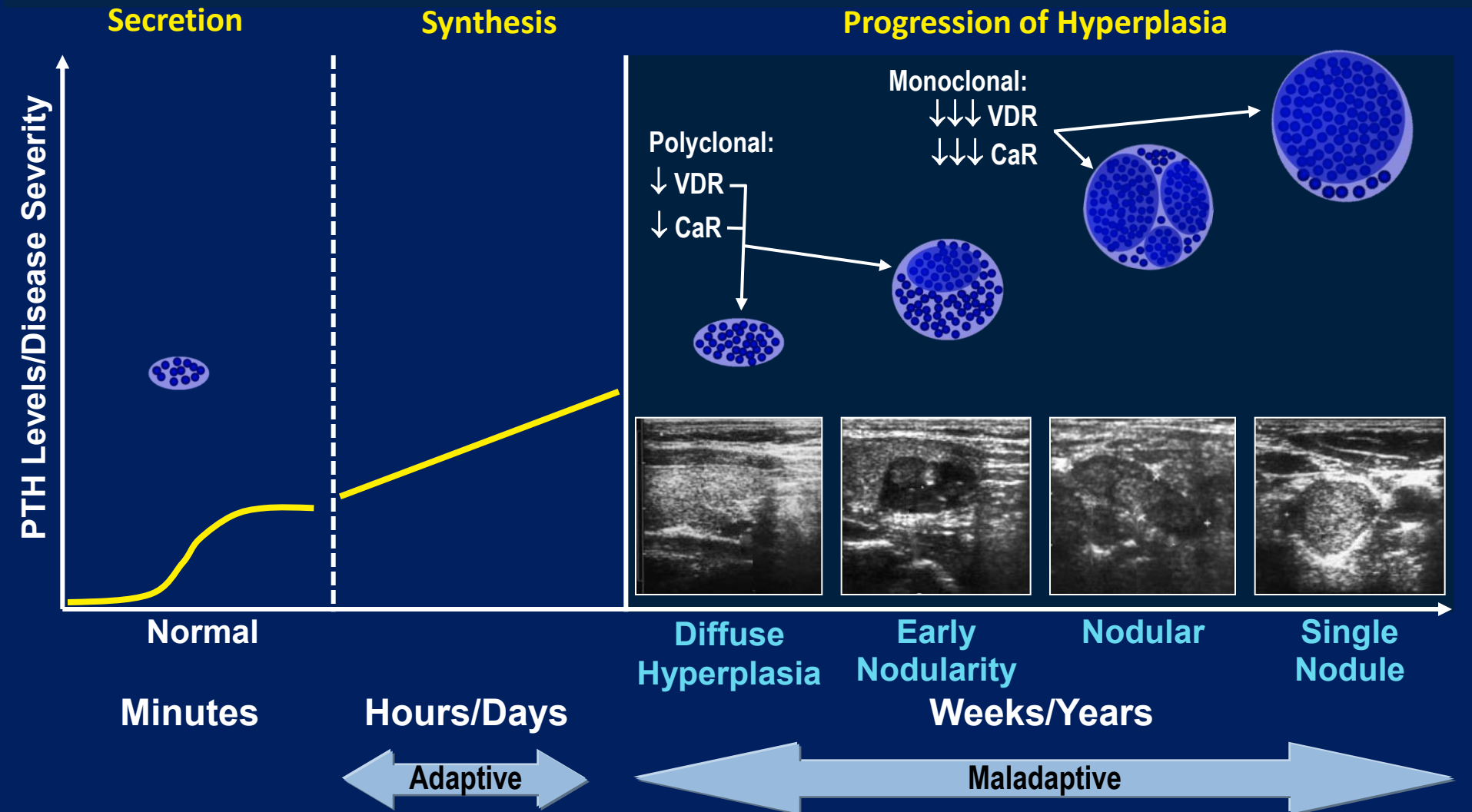


Figure 1. Axis of PTH, FGF-23, and 1,25-dihydroxyvitamin D. PTH, FGF-23, and 1,25(OH)₂D interact and regulate one another via classic negative endocrine feedback loops that affect calcium and phosphate transport in the kidney, bone, and intestine. Created with Biorender. Abbreviations: 1,25D, 1,25-dihydroxyvitamin D; FGF-23, fibroblast growth factor-23; PTH, parathyroid hormone.



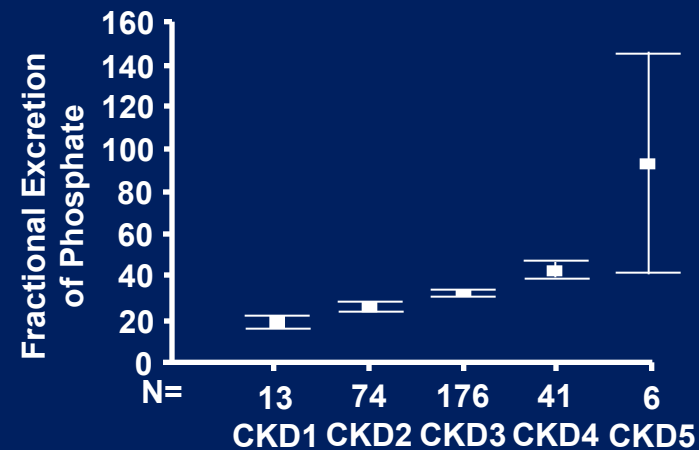
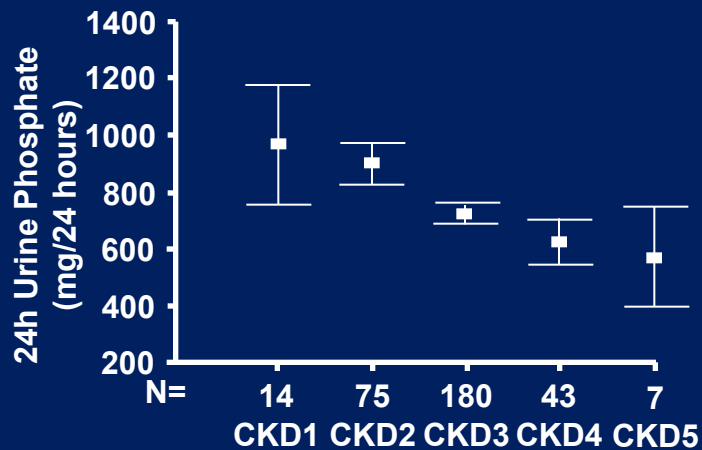
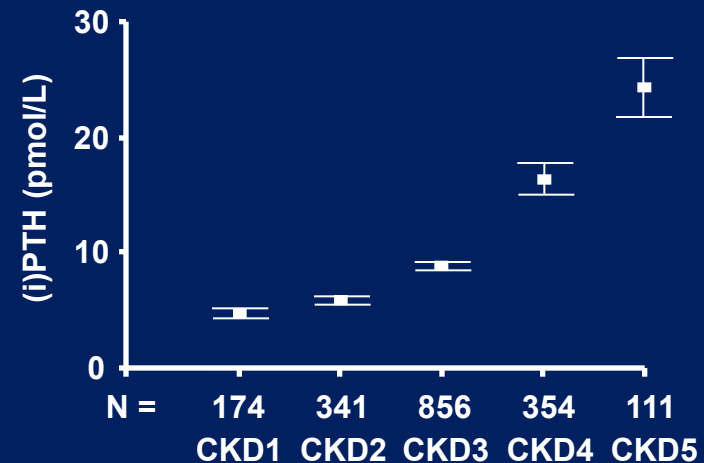
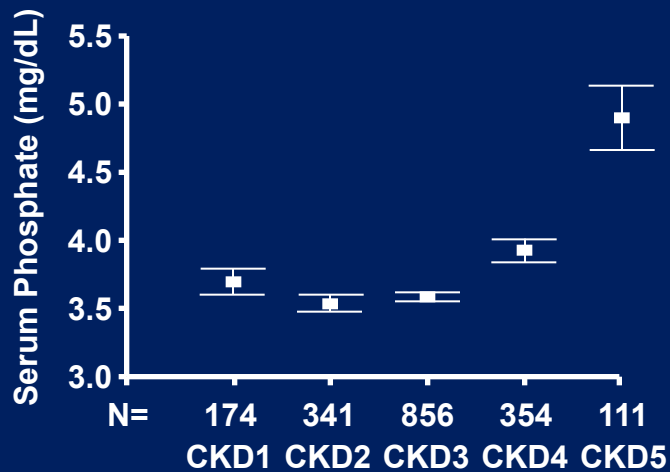
Progression of Parathyroid Gland Hyperplasia in Secondary HPT



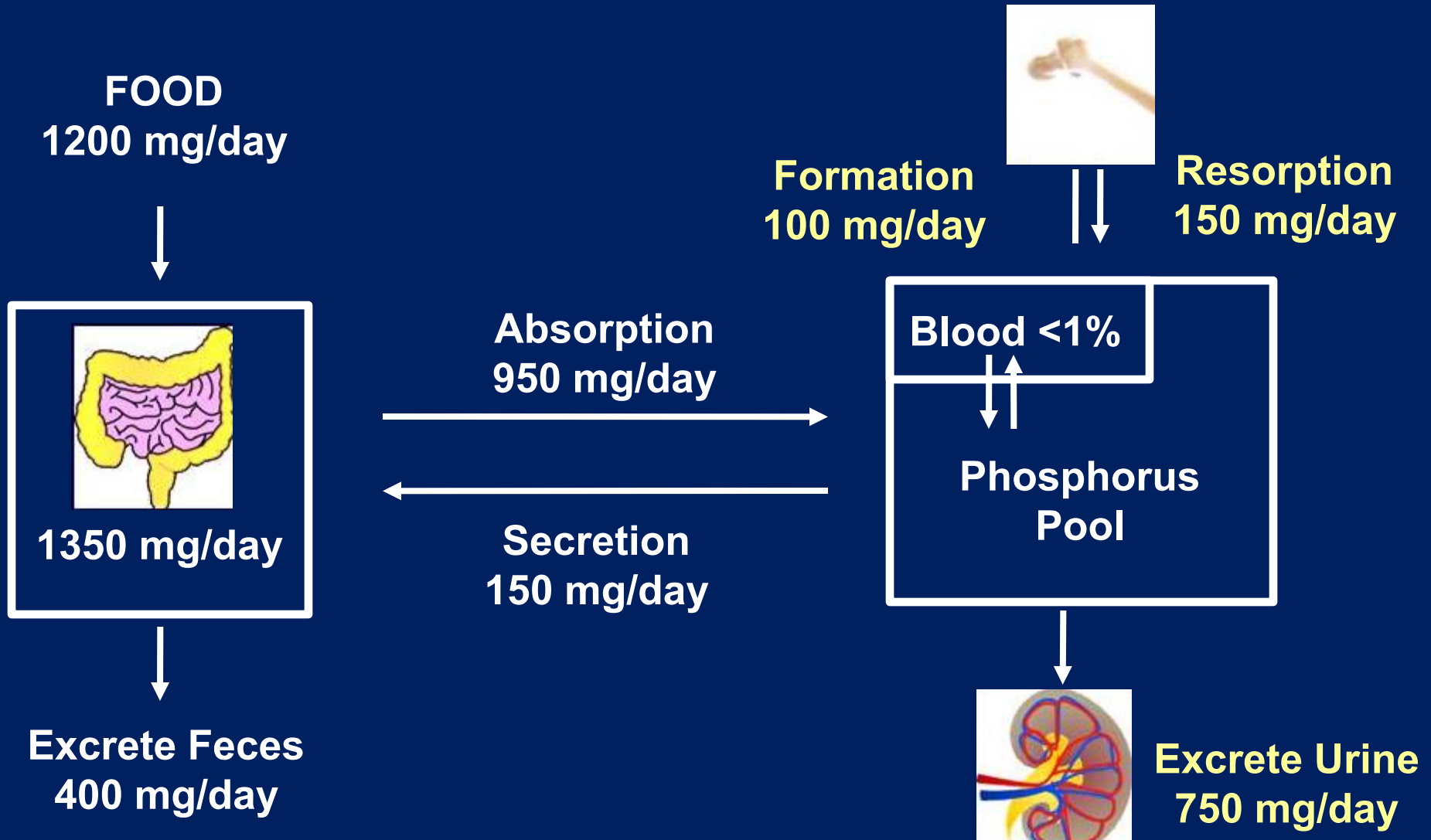
VDR = vitamin D receptor; CaR = calcium-sensing receptor.

Adapted from Rodriguez M, et al. *Am J Physiol Renal Physiol*. 2005;288:F253-F264; Adapted from Tominaga Y, et al. *Semin Surg Oncol*. 1997;13:78-86; Adapted from Pavlovic D, et al. *Sci World J* 2006;6:1599-1608.

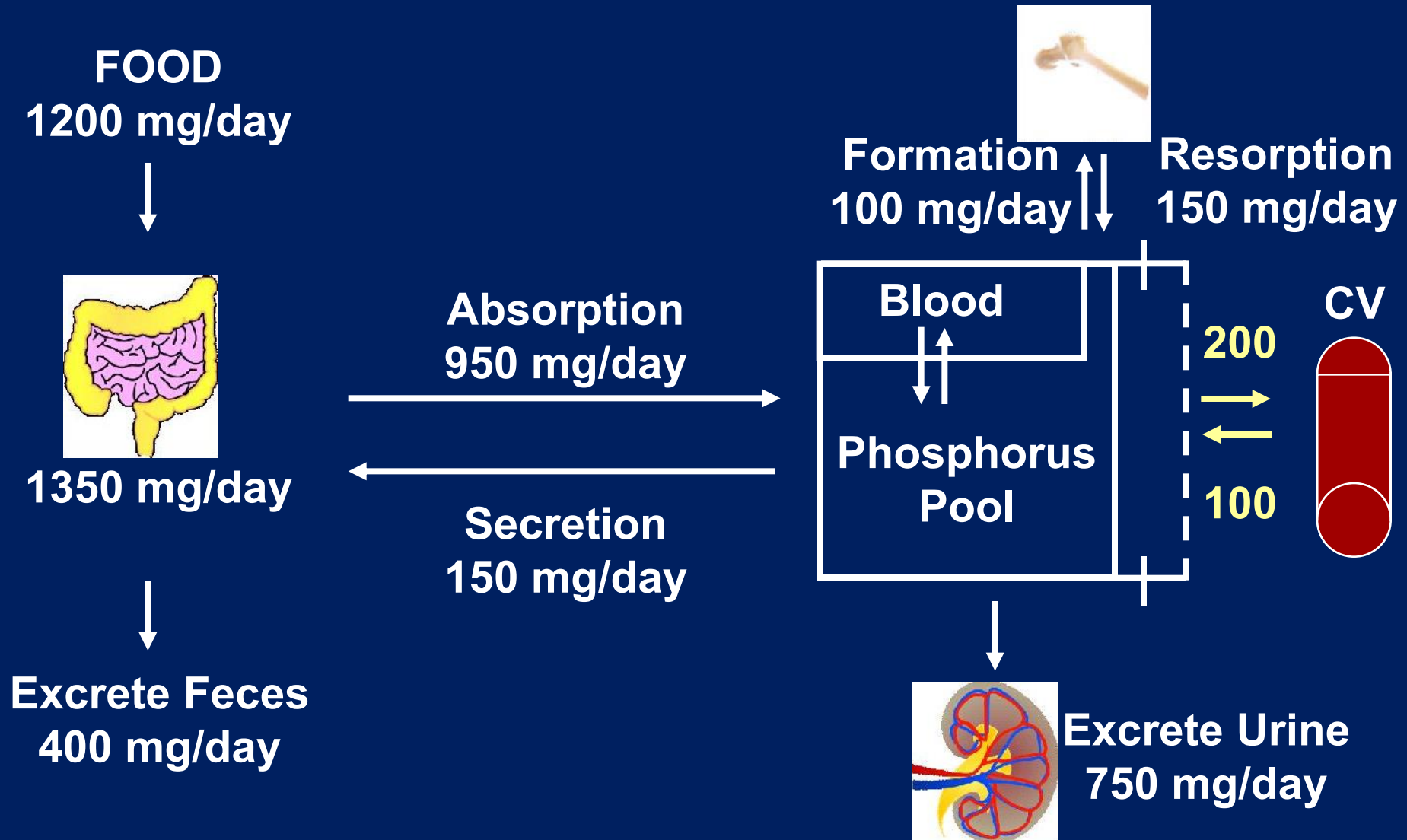
Phosphate Increases With Decreased Kidney Function



Phosphorus Balance Is Lost in CKD

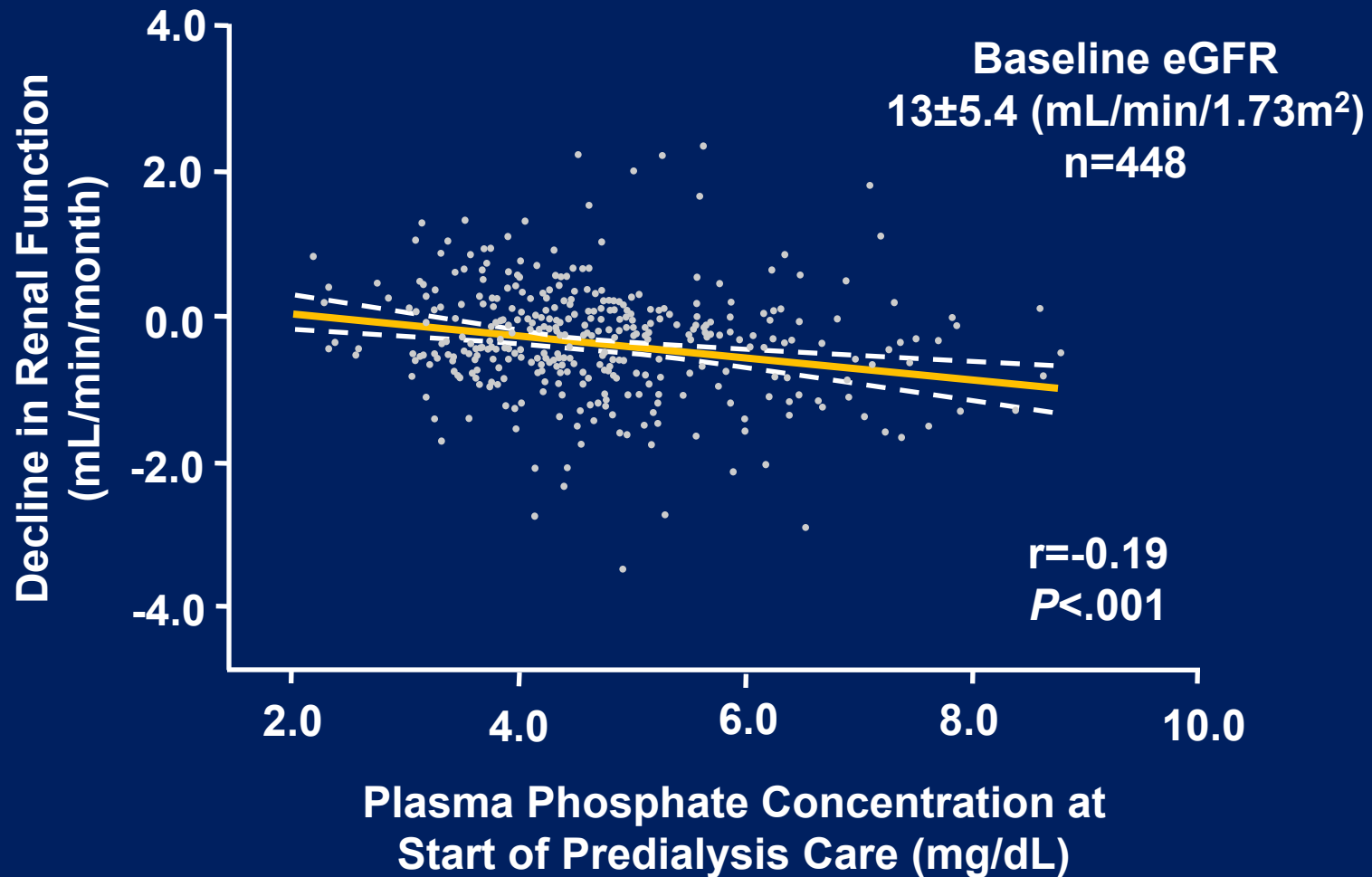


Phosphorus Balance Is Lost in CKD

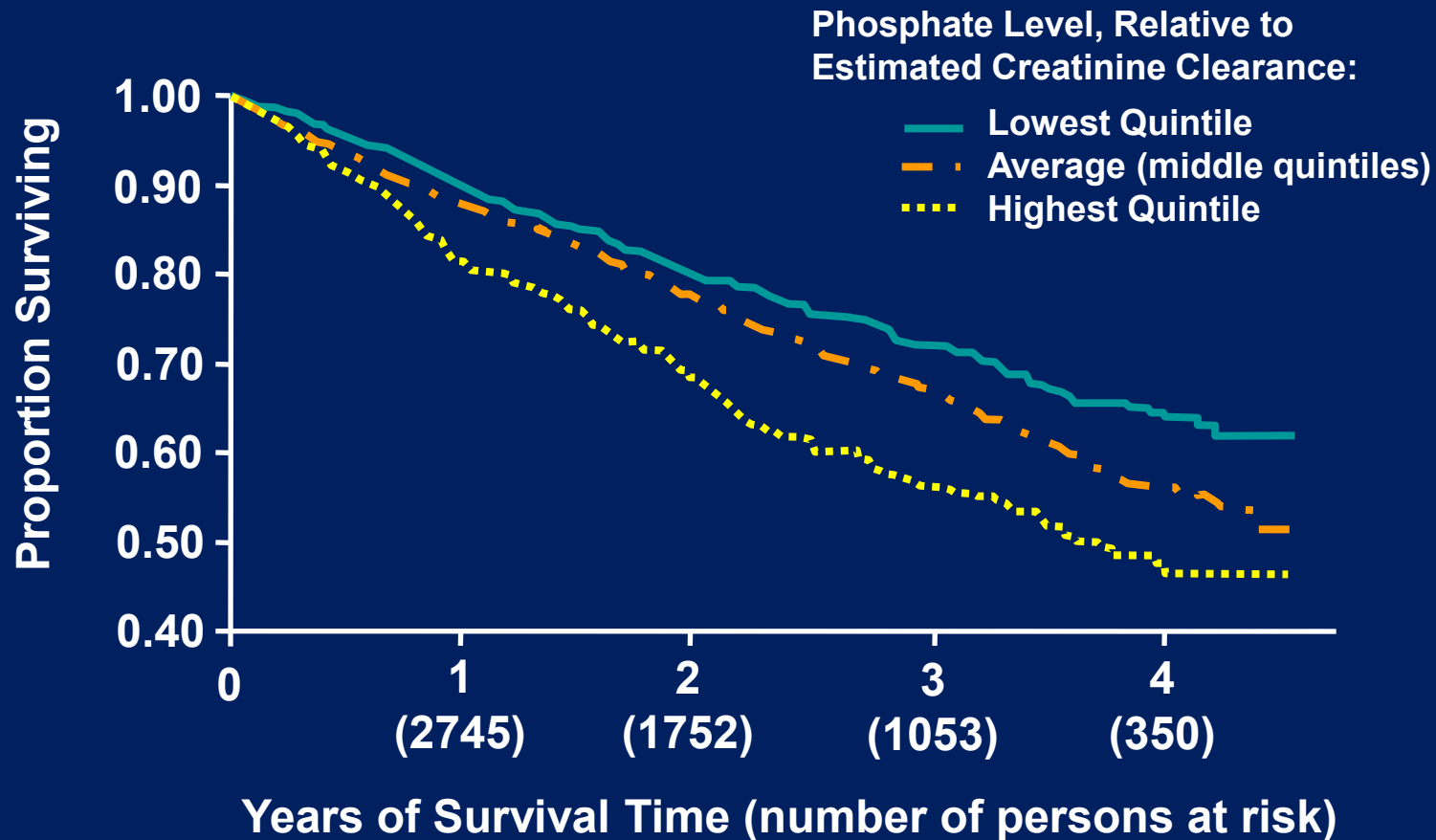


Potential Deleterious Effects of Disorders of Mineral Metabolism

Higher Serum Phosphorus Is Associated With More Rapid Decline in Renal Function

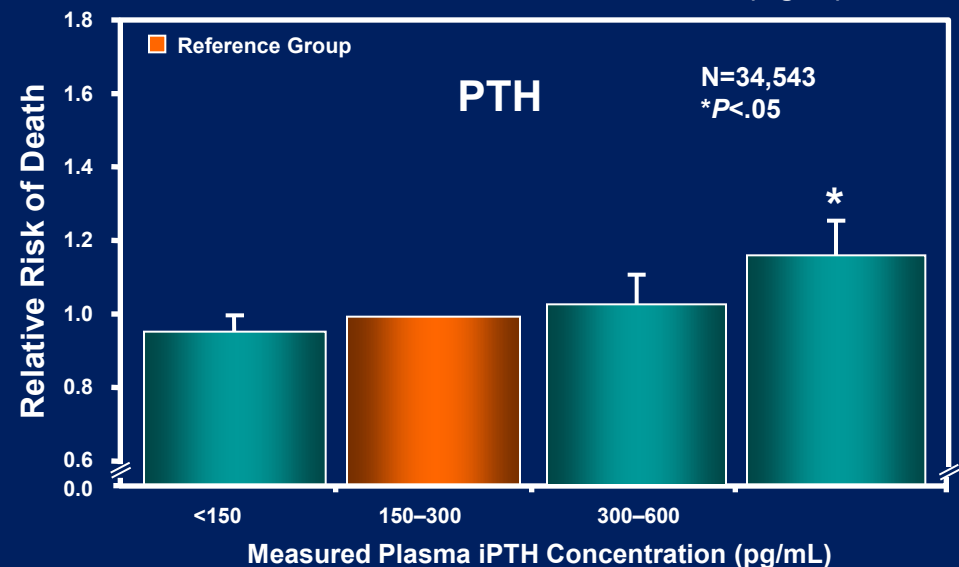
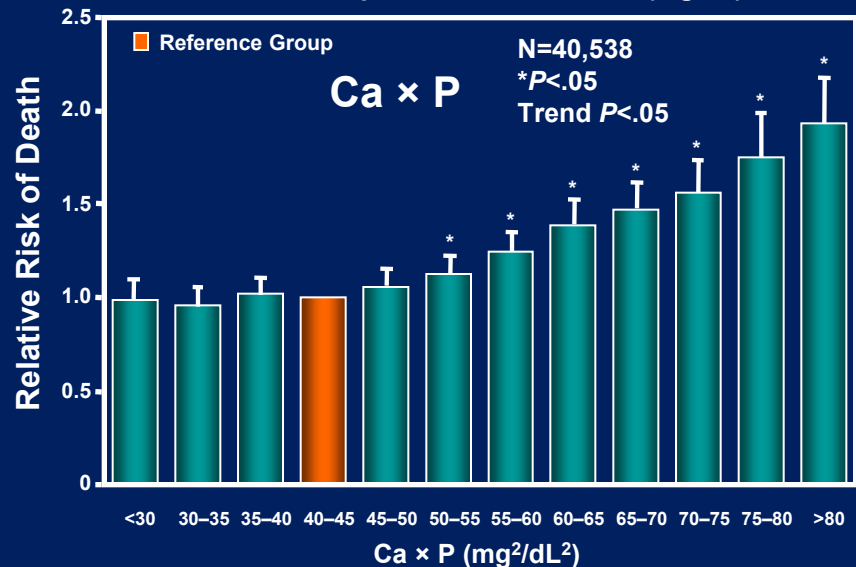
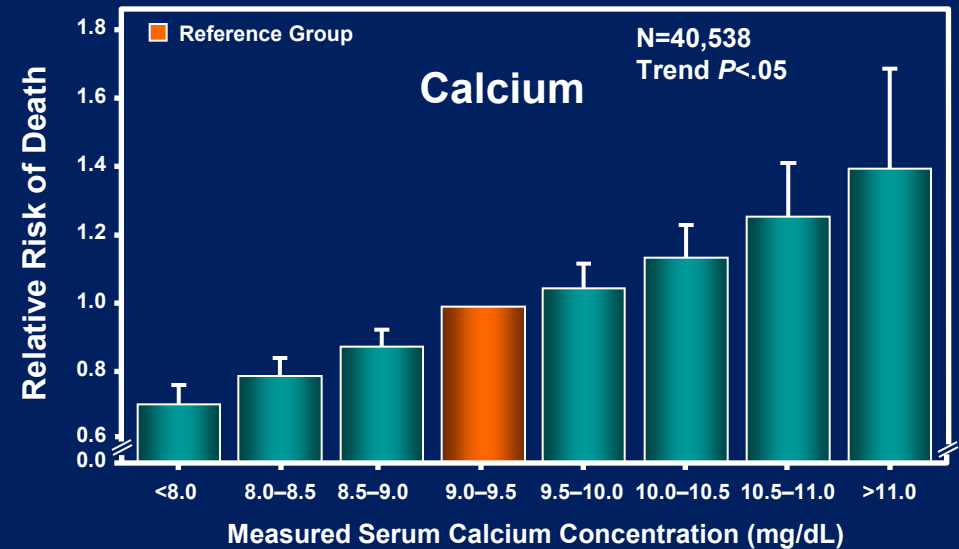
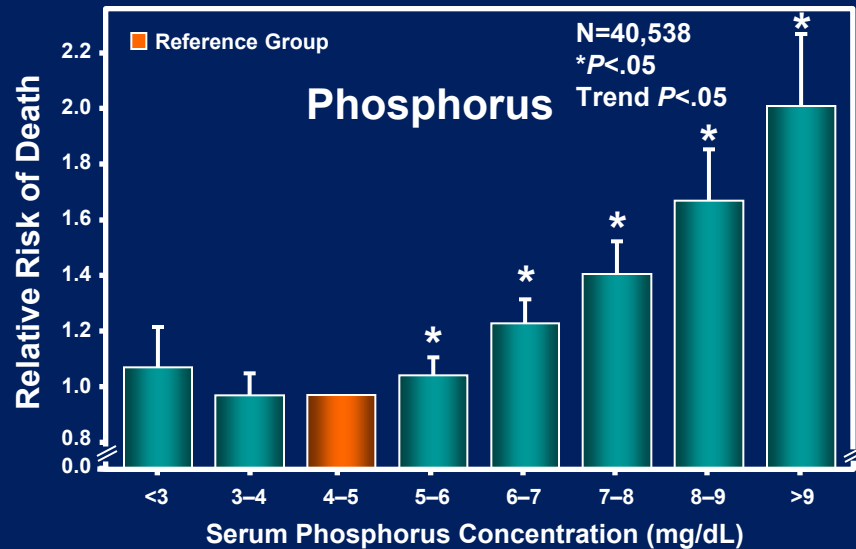


Serum Phosphorus and Mortality

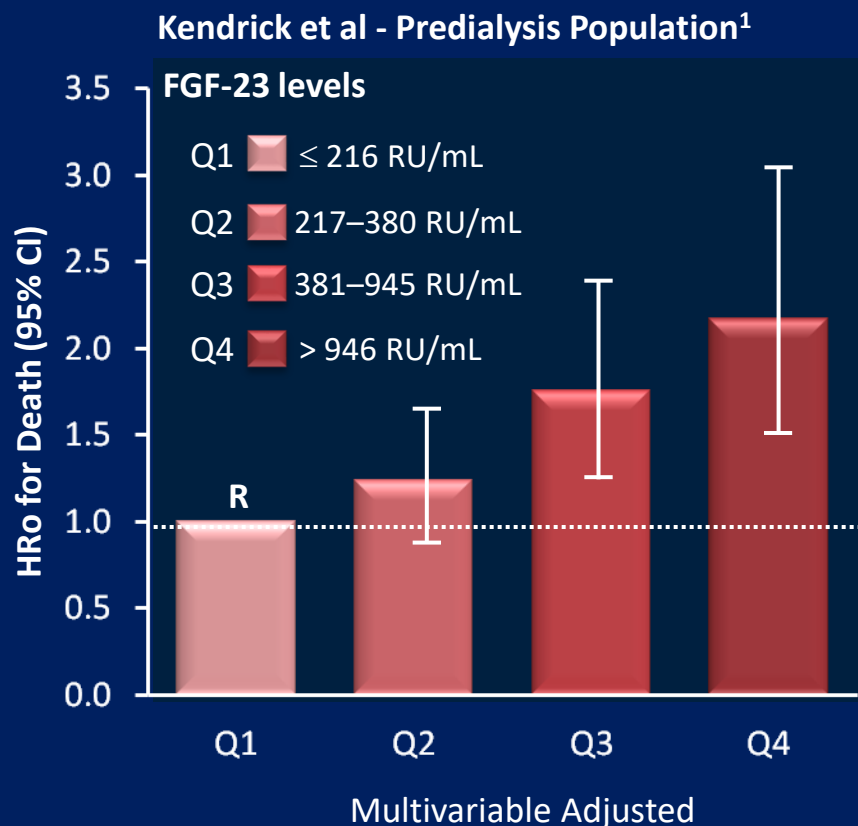


- Estimates of survival significantly different between patients with relative phosphorus levels in highest and lowest quintiles predicted by estimated creatinine clearance ($P < .001$)

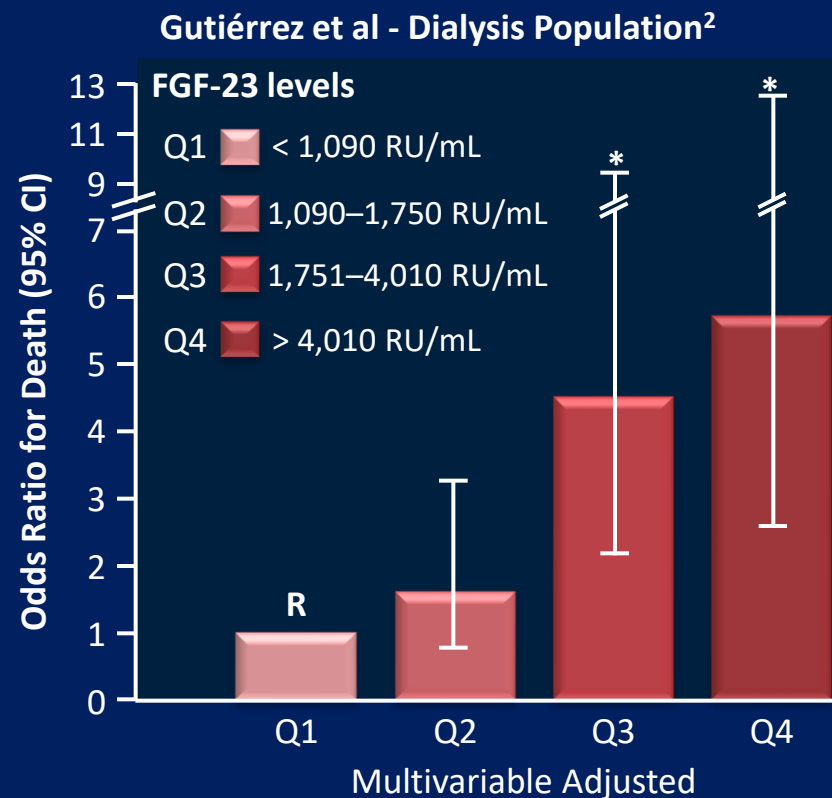
Observational Data Suggest a Link Between Laboratory Measures and Mortality



Across the Spectrum of CKD, Elevated FGF-23 Levels Were Associated With Higher Mortality Risk



N = 1,099



N = 400

* $P < 0.05$

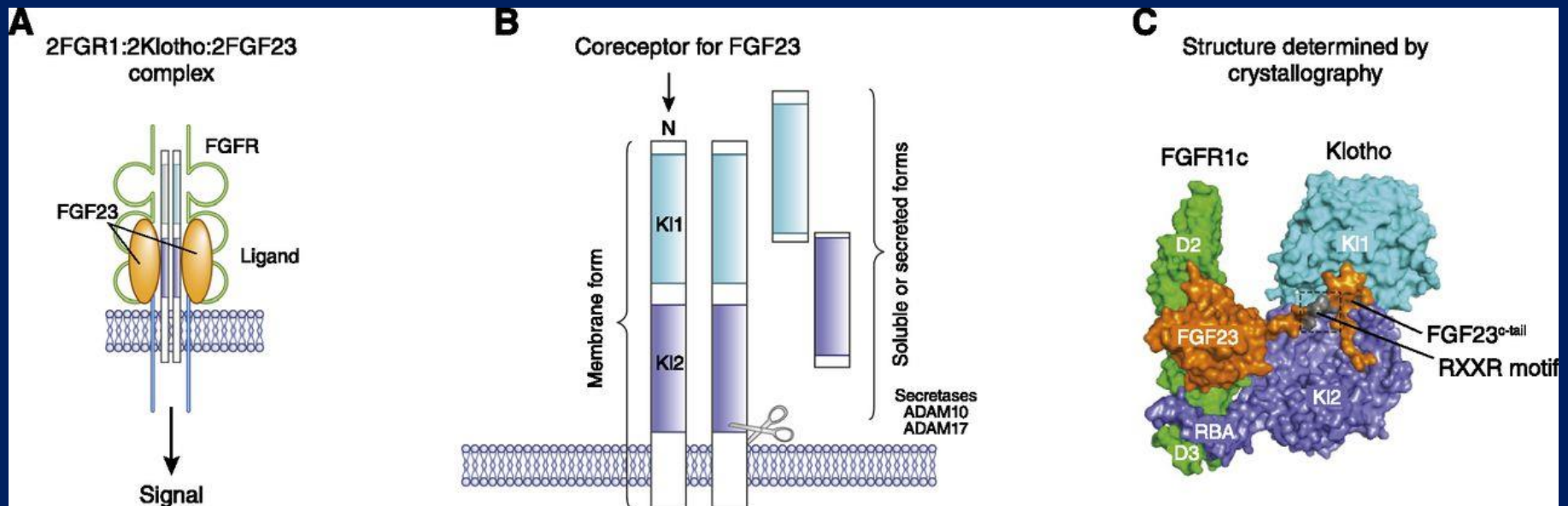
Patients in the highest quartiles of FGF-23 levels had the highest risk of mortality as compared with subjects in the lowest quartiles^{1,2}

Q = quartile; R = reference.

1. Kendrick J, et al. *J Am Soc Nephrol*. 2011;22:1913-1922.

2. Gutiérrez OM, et al. *N Engl J Med*. 2008;359:584-592.

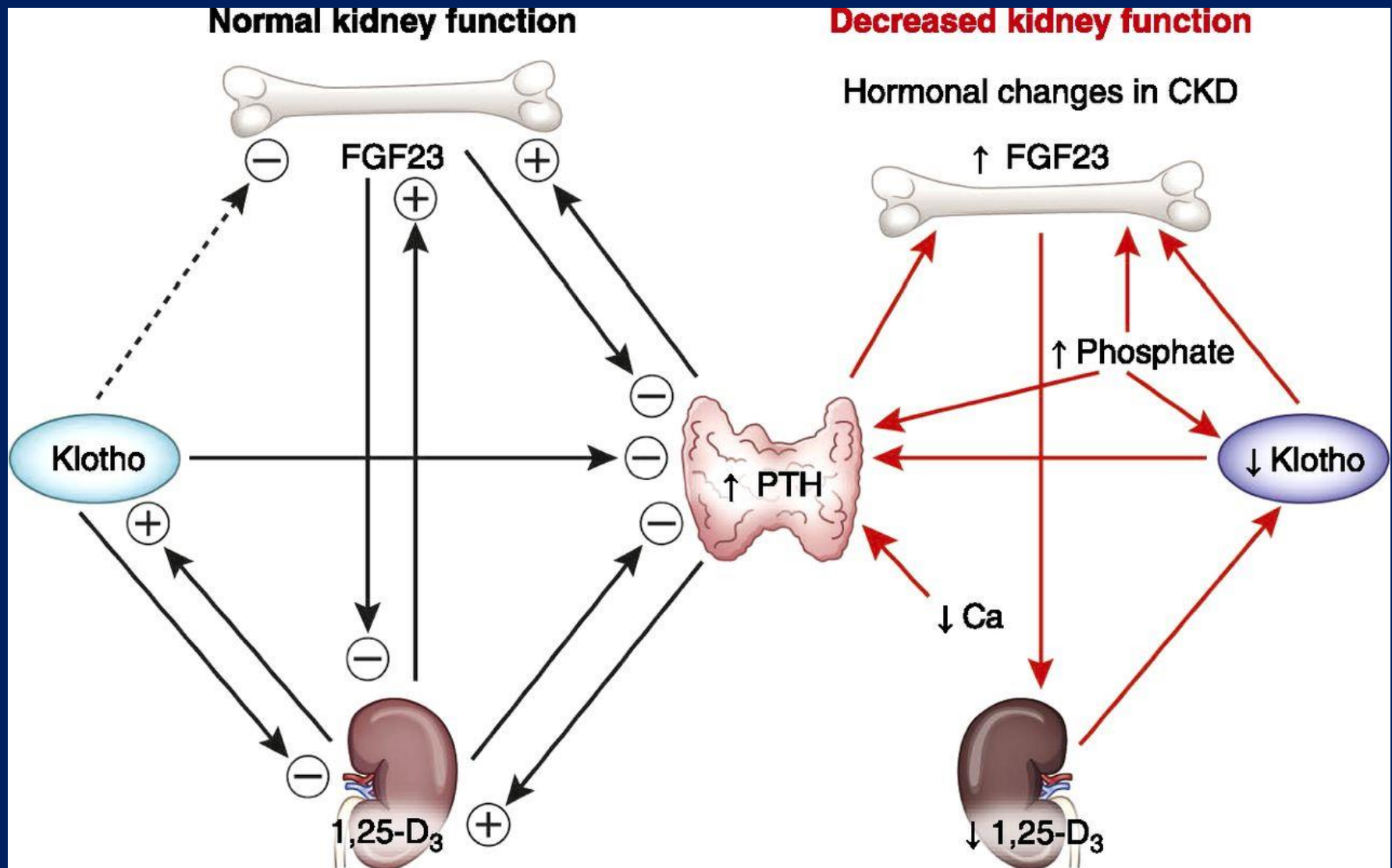
Overview of Klotho protein.



Klotho—the fountain of youth?

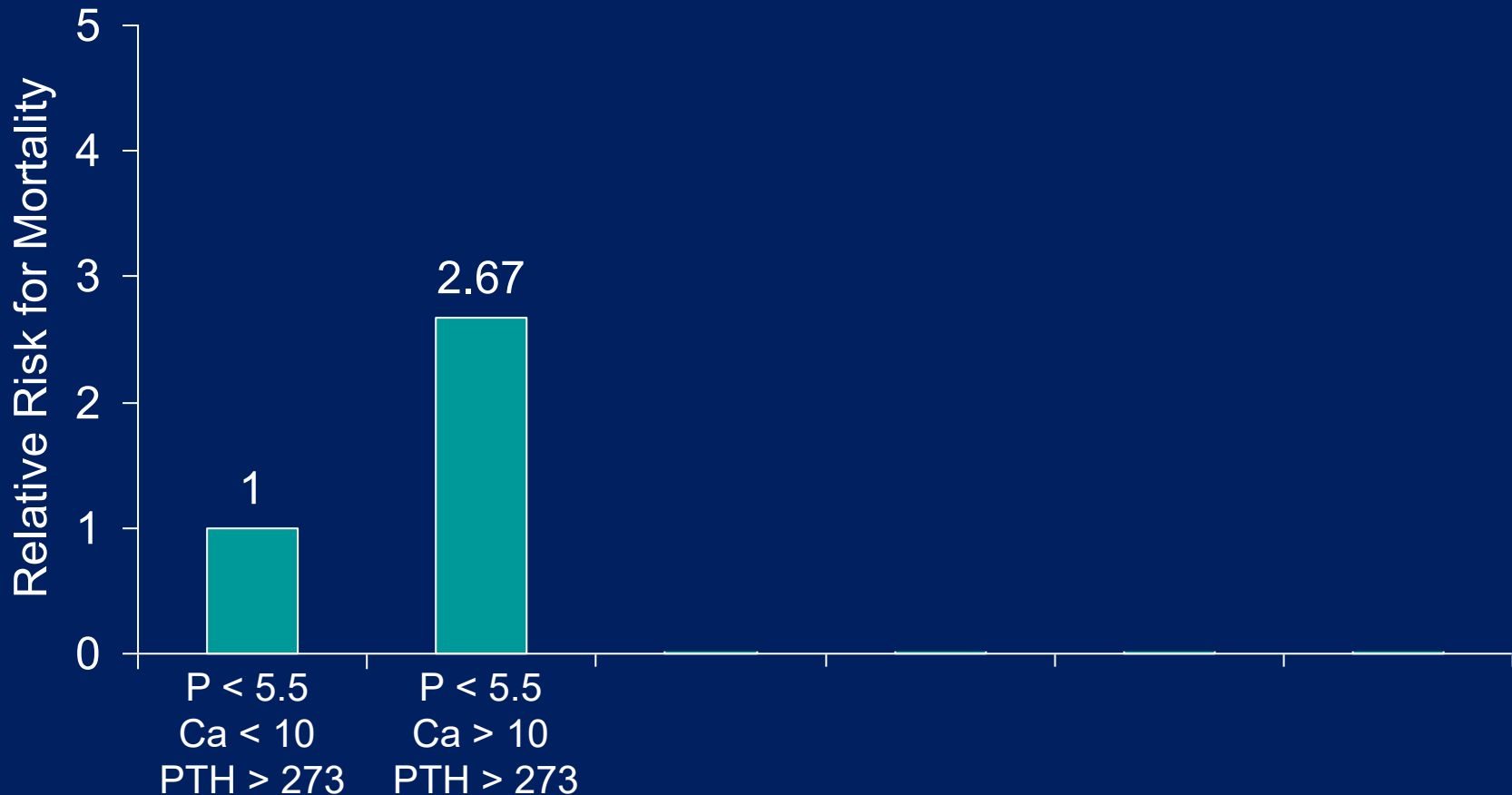
- Protects against renal fibrosis and promotes kidney regeneration after ischemic injury
- Inhibits vascular calcification.
- Reduces oxidative stress by inhibition of insulin and insulin-like growth factor-1 signaling pathways.
- Increases nitric oxide production /improves vasodilatation.
- Inhibits cell senescence and apoptosis
- Preserves stem cells

Physiologic and pathophysiologic role of Klotho in mineral metabolism with preserved and decreased kidney function.

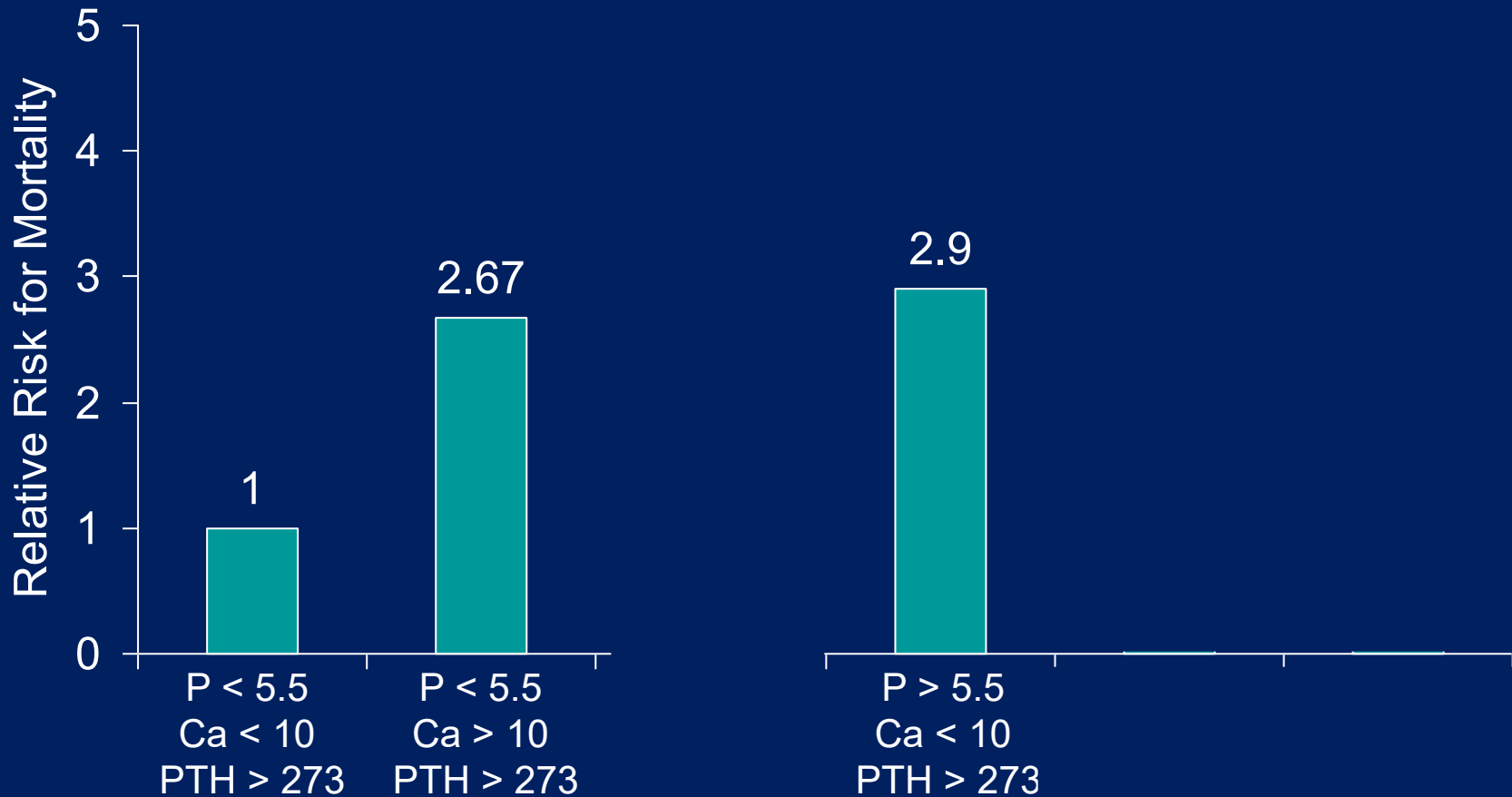


Calcification

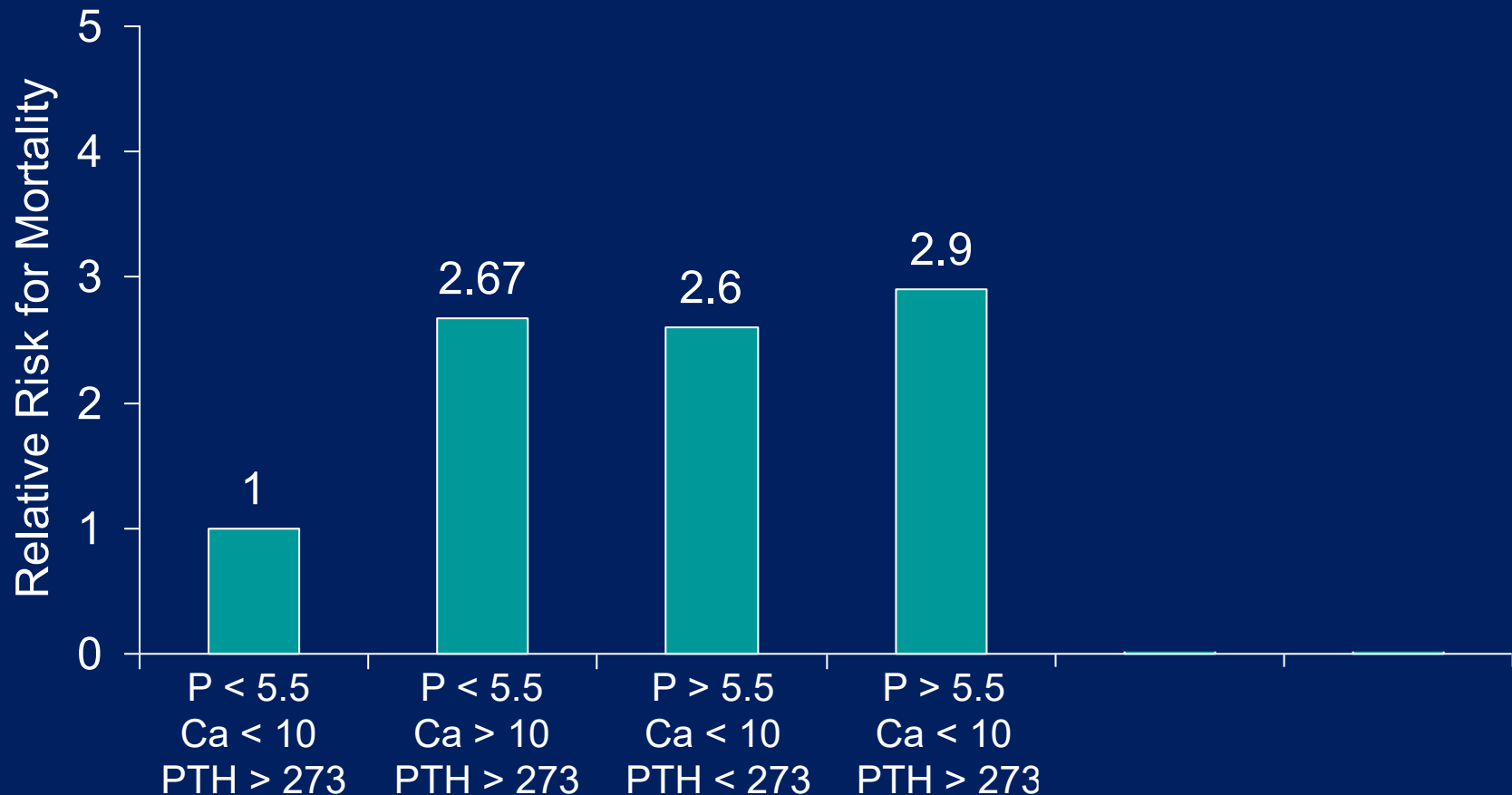
Disordered Mineral Metabolism and Mortality: A Complex Relationship



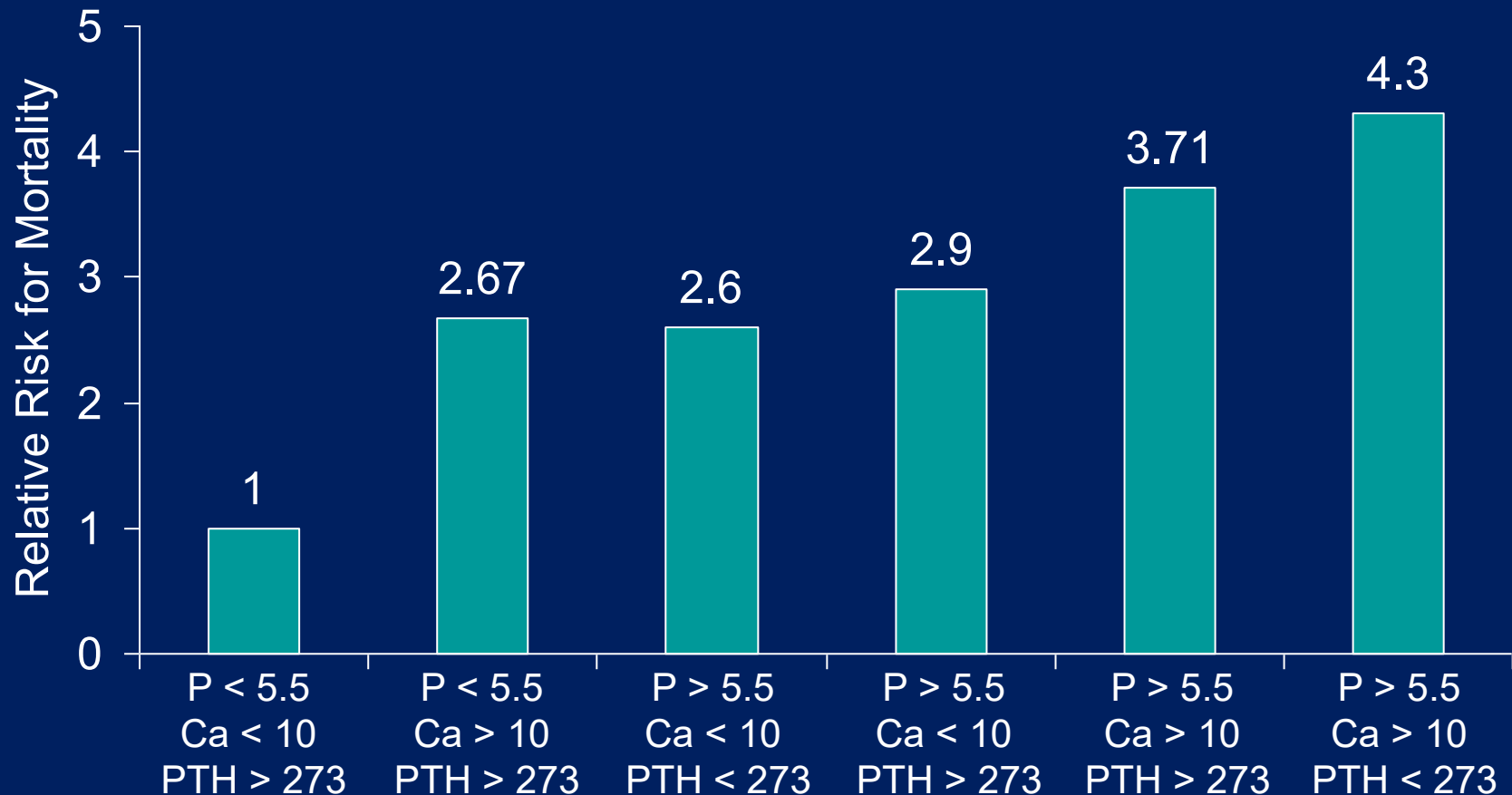
Disordered Mineral Metabolism and Mortality: A Complex Relationship



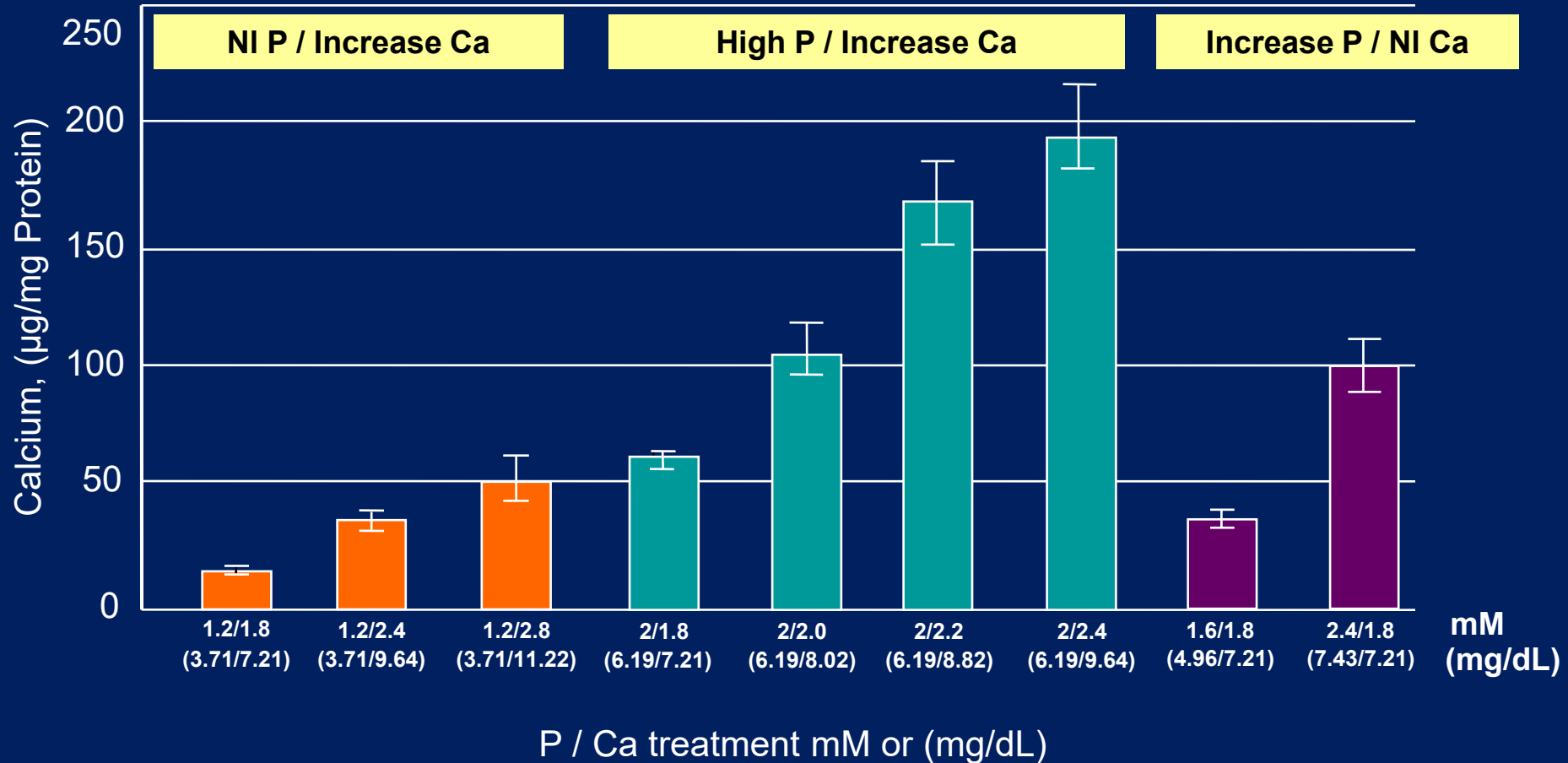
Disordered Mineral Metabolism and Mortality: A Complex Relationship

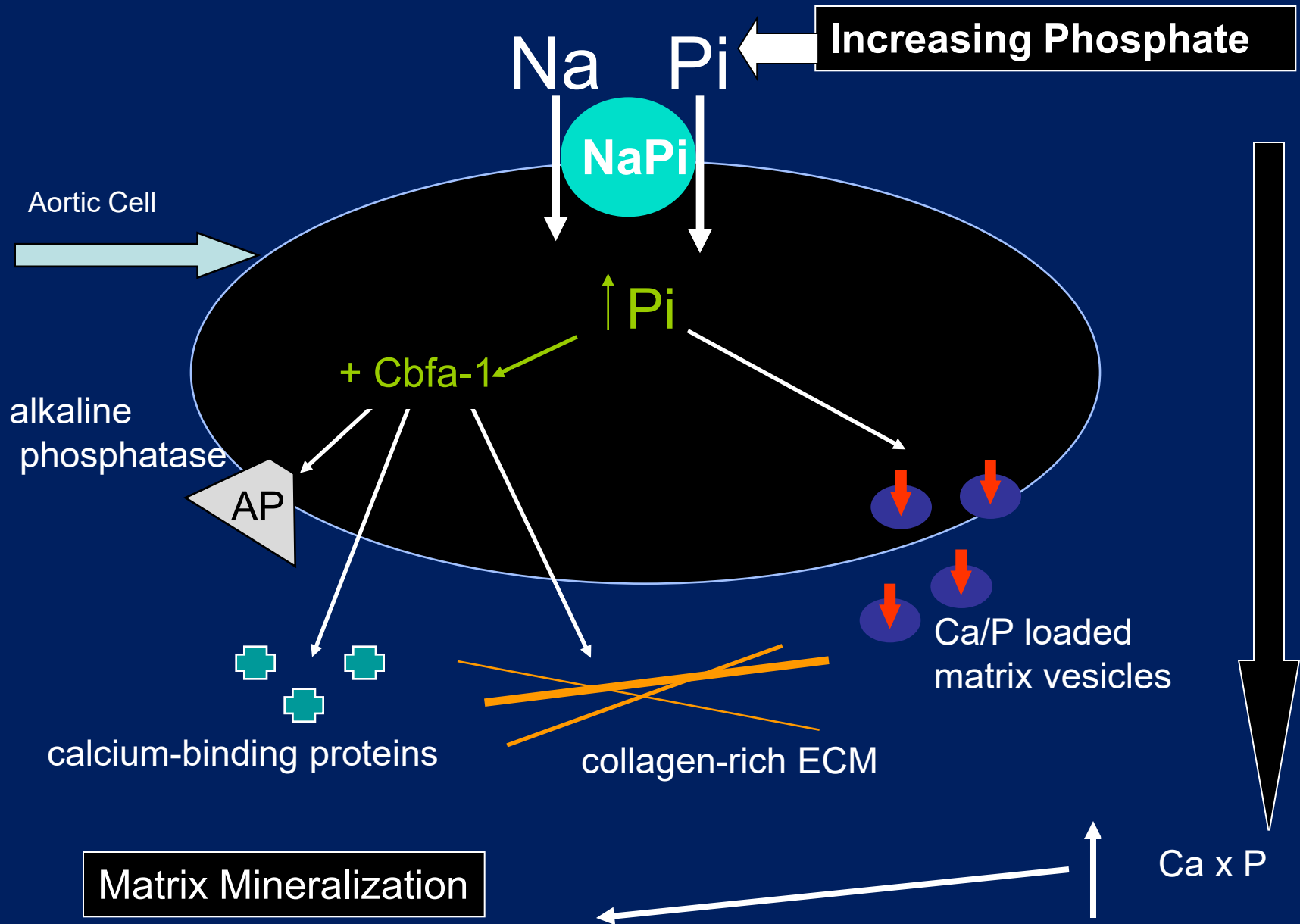


Disordered Mineral Metabolism and Mortality: A Complex Relationship

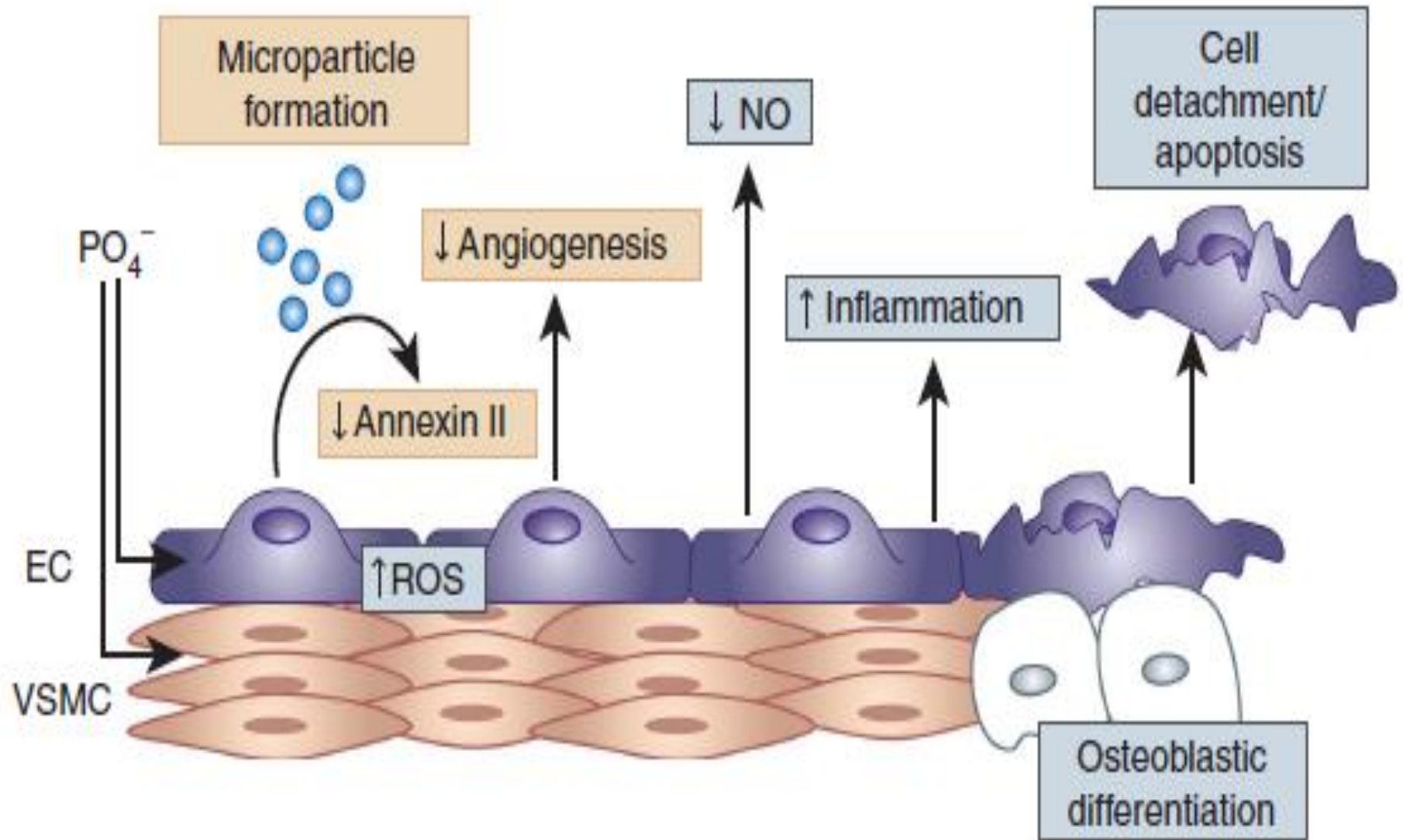


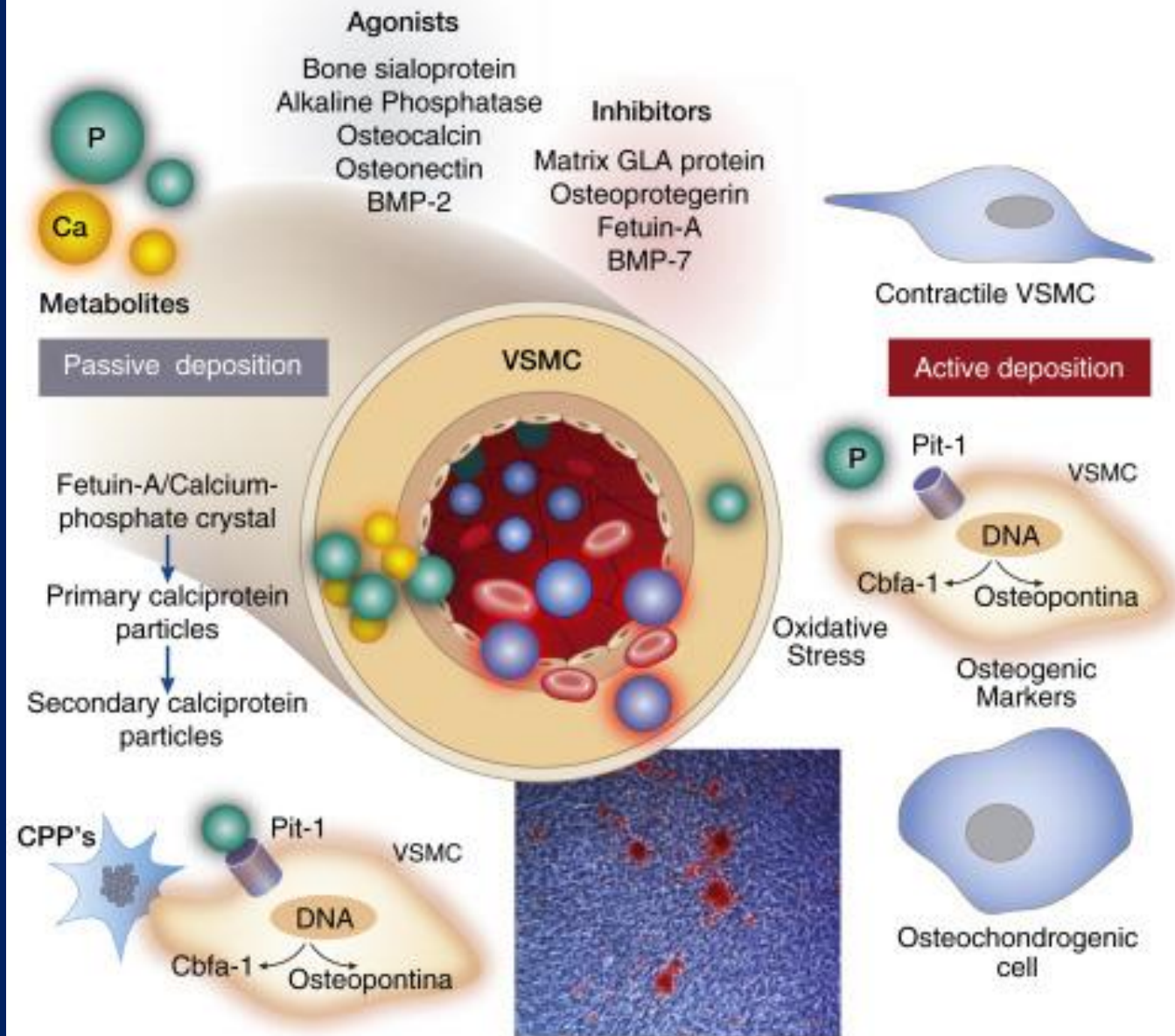
Ca & P Induce Calcification in SMC





Effects of Phosphate on Vascular Health

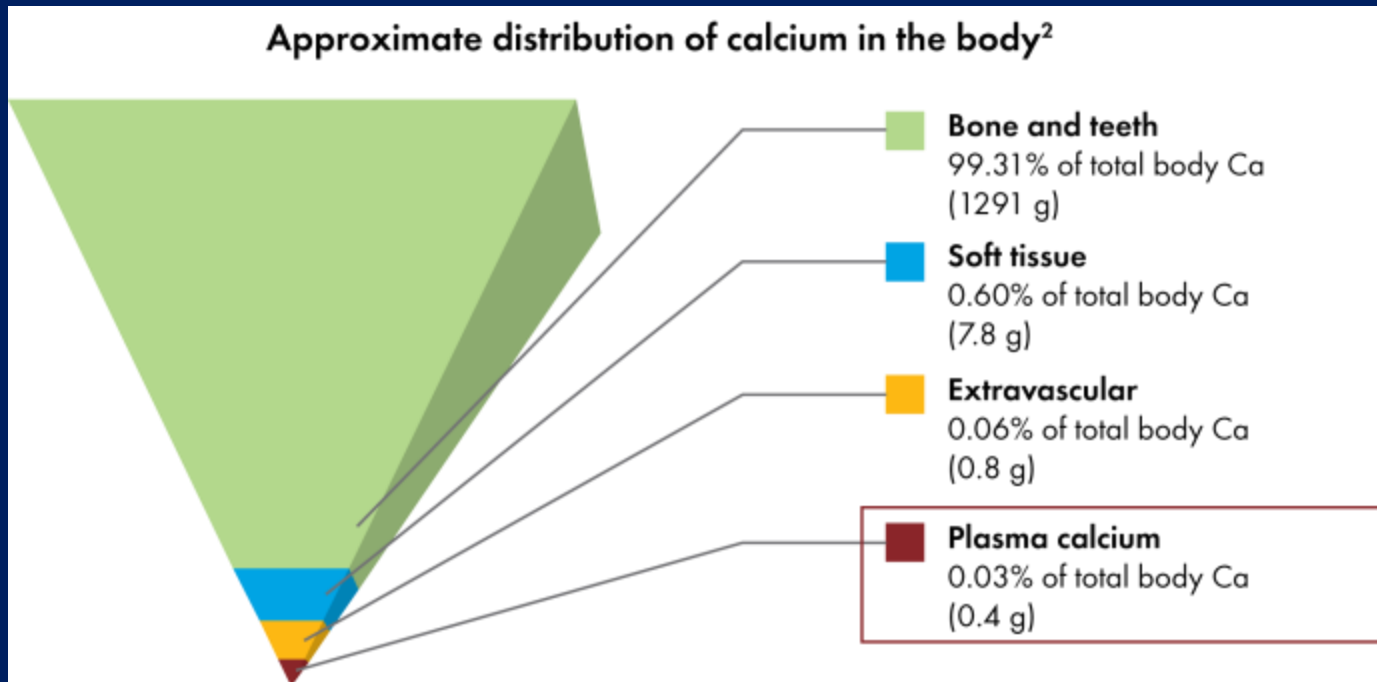




Vascular and Valvular Calcification:

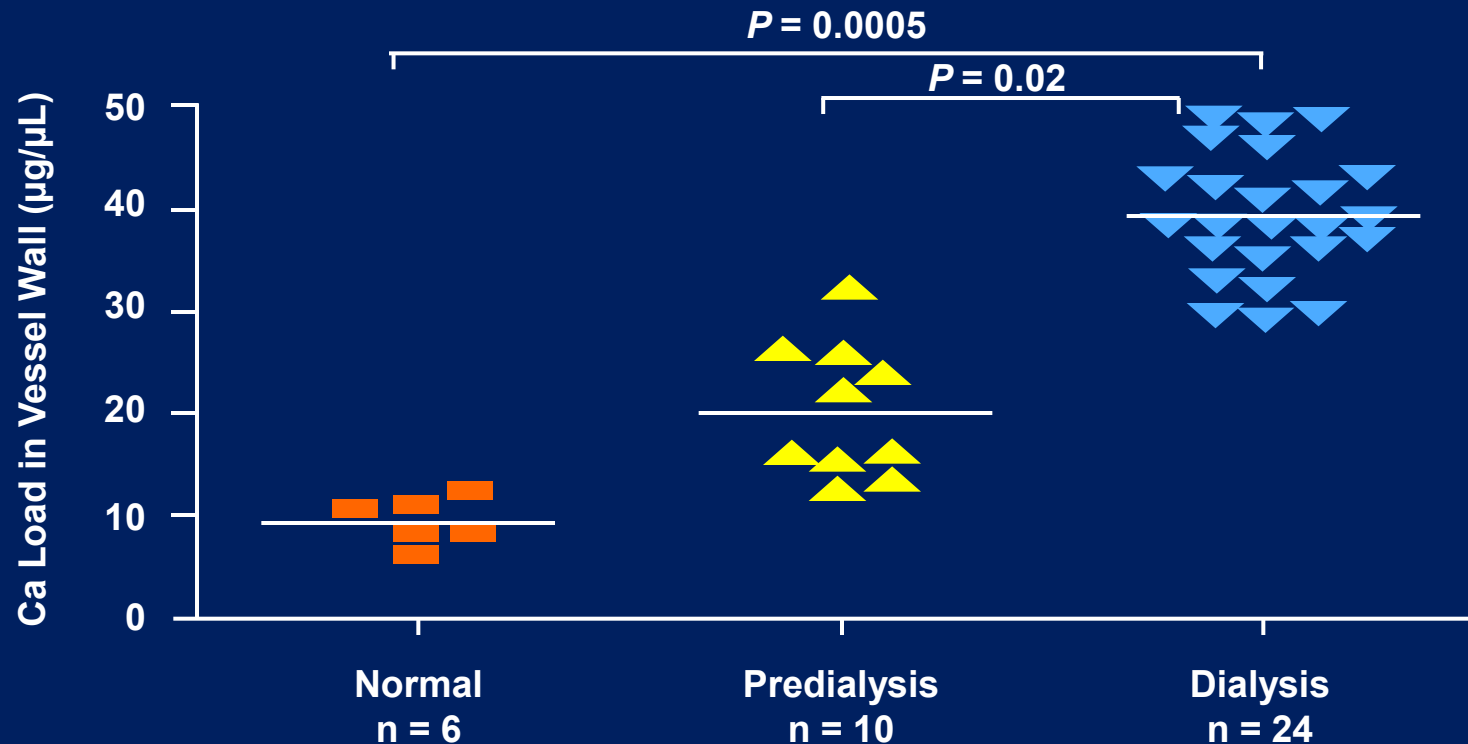
*Potential Consequences of Disorders of
Divalent Ion Metabolism*

Calcium distribution in the body

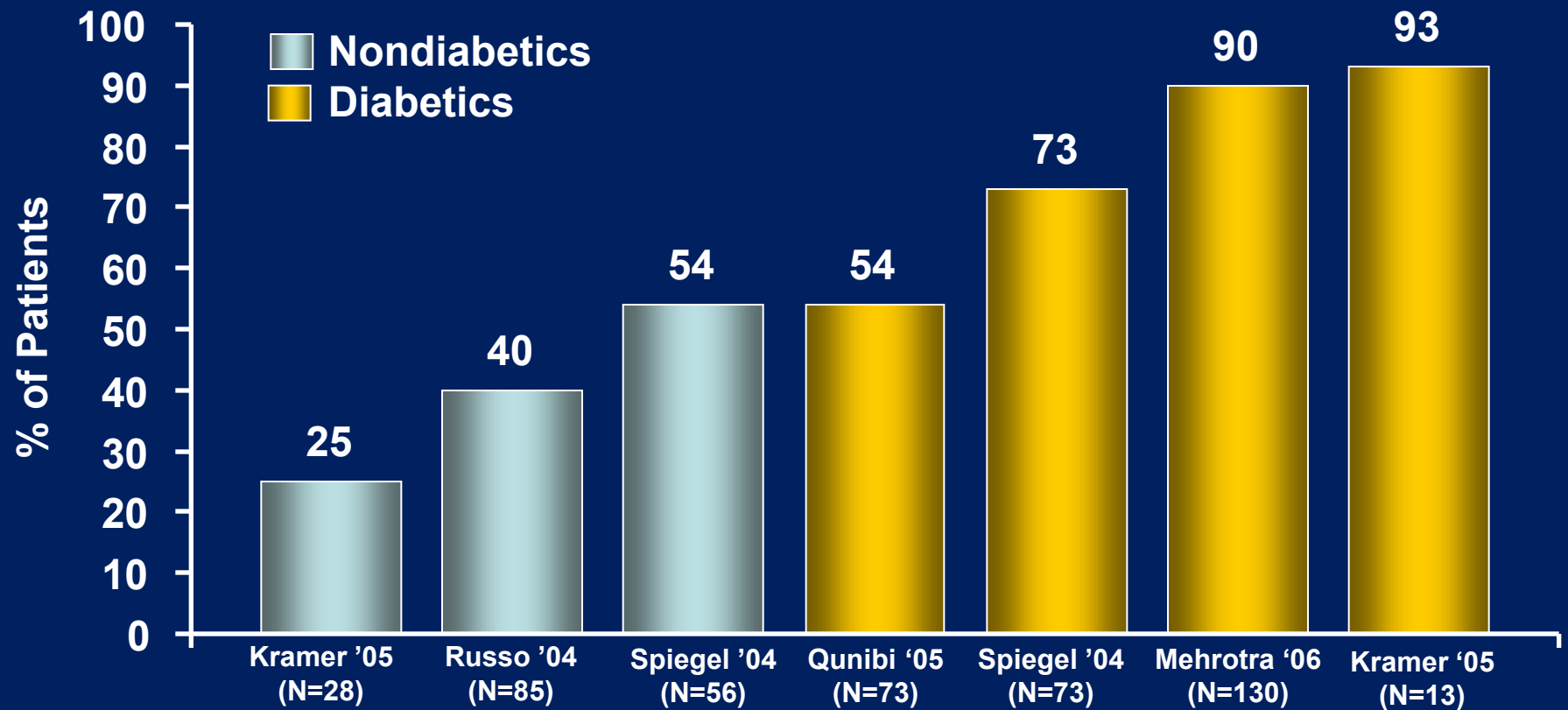


References: 1. Houillier P, Froissart M, Maruani G, Blanchard A. What serum calcium can tell us and what it can't. *Nephrol Dial Transplantation*. 2006;21:29-32. 2. Nordin BEC, ed. *Calcium, Phosphate and Magnesium Metabolism: Clinical Physiology and Diagnostic Procedures*. New York, NY: Churchill Livingstone; 1976. 3. Hosking DJ, Chamberlain MJ. Calcium balance in chronic renal failure: a study using in vivo neutron activation analysis. *Q J Med*. 1973;42:467-479. 4. Bushinsky DA. Contribution of intestine, bone, kidney, and dialysis to extracellular fluid calcium content. *Clin J Am Soc Nephrol*. 2010;5(suppl 1):S12-S22.

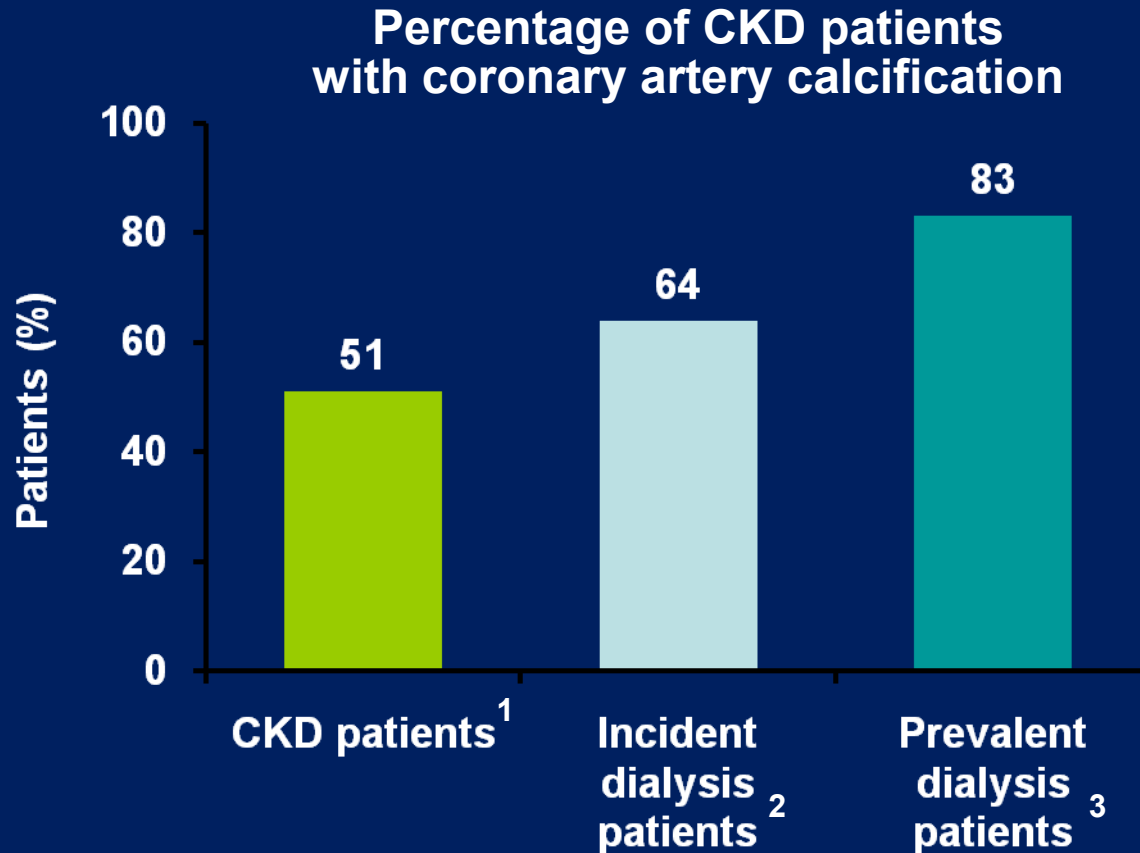
Vascular Calcium Loading in Children With CKD



Prevalence of Coronary Calcification in CKD Patients Not on Dialysis



Across CKD Patient Populations, Calcification is a Common, Progressive Consequence of the Disease

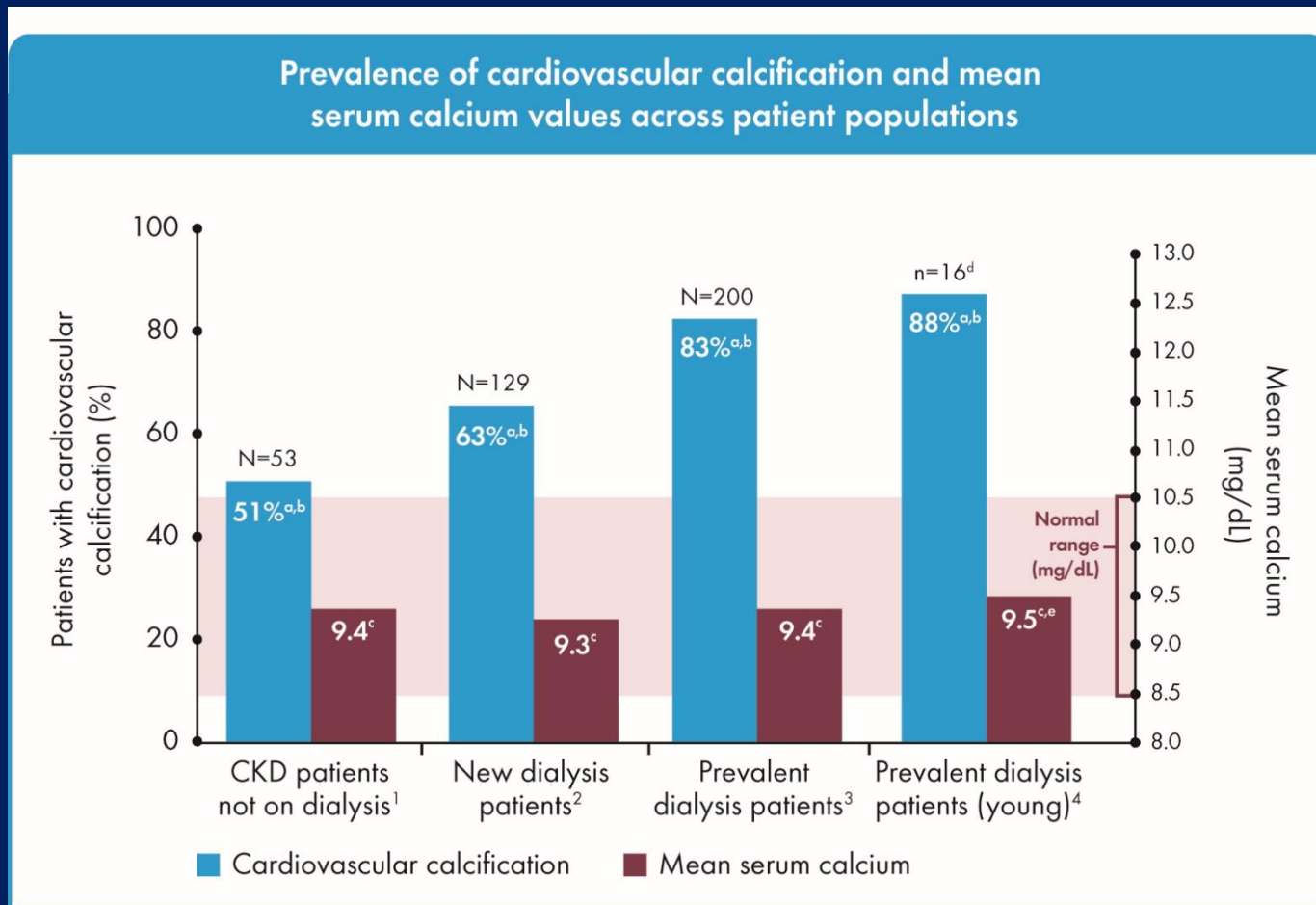


1. Russo D, Corrao S, Miranda I, et al. *Am J Nephrol*. 2007;27:152-158.

2. Spiegel DM, Raggi P, Mehta R, et al. *Hemodialysis Int*. 2004;8:265-272.

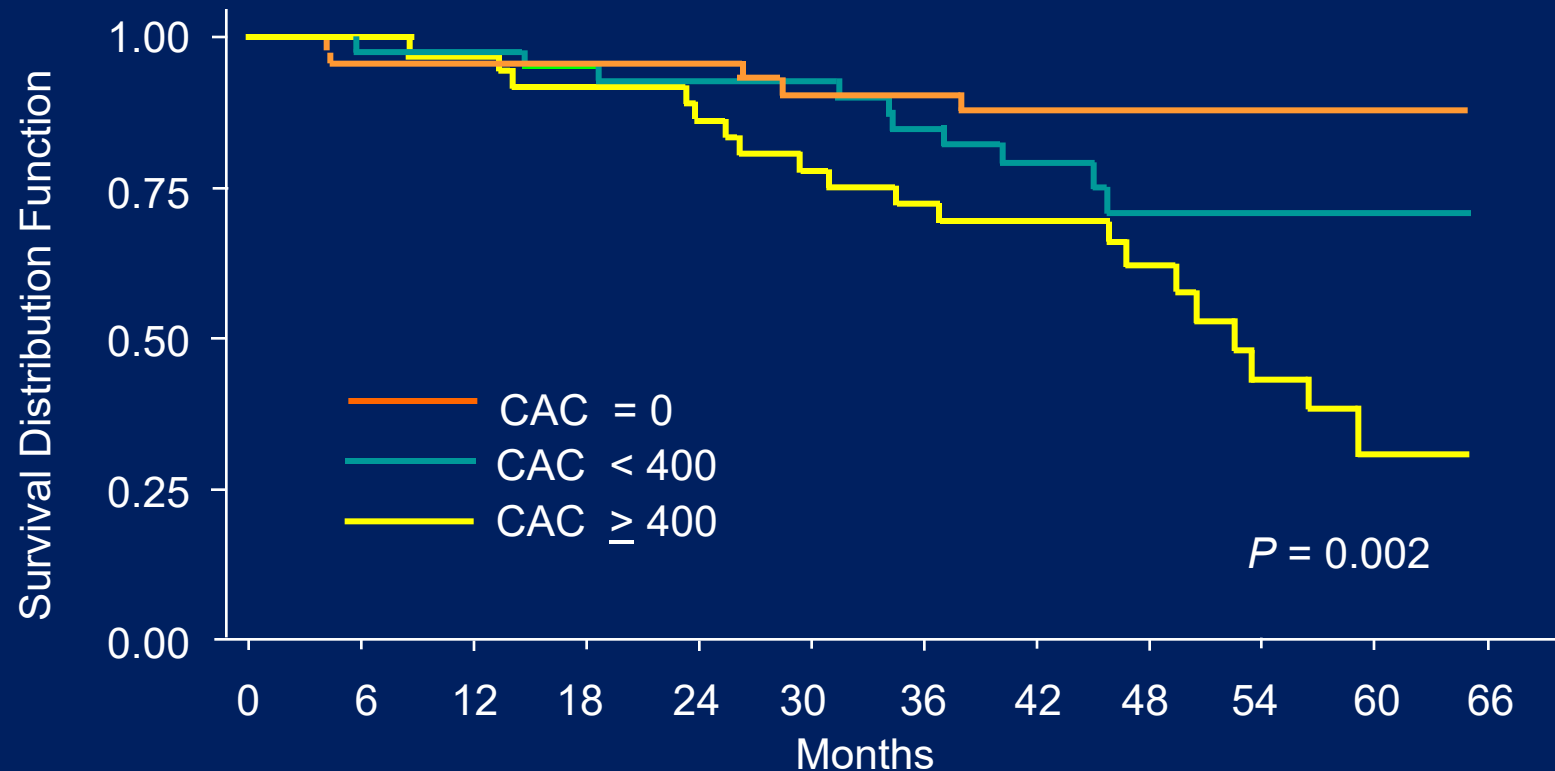
3. Chertow GM, Burke SK, Raggi P; *Kidney Int*. 2002;62:245-252.

Calcification is independent of serum calcium levels ¹⁻⁵



1. Russo D, et al. *Am J Nephrol.* 2007;27:152-158. 2. Block GA, et al. *Kidney Int.* 2005;68:1815-1824. 3. Chertow GM, et al. *Kidney Int.* 2002;62:245-252. 4. Goodman WG, et al. *N Engl J Med.* 2000;342:1478-1483. 5. Kidney Disease: Improving Global Outcomes (KDIGO) CKD-MBD Work Group. KDIGO *Kidney Int.* 2009;76(suppl 113):S1-S130.

Coronary Artery Calcification Is Associated With Increased Mortality

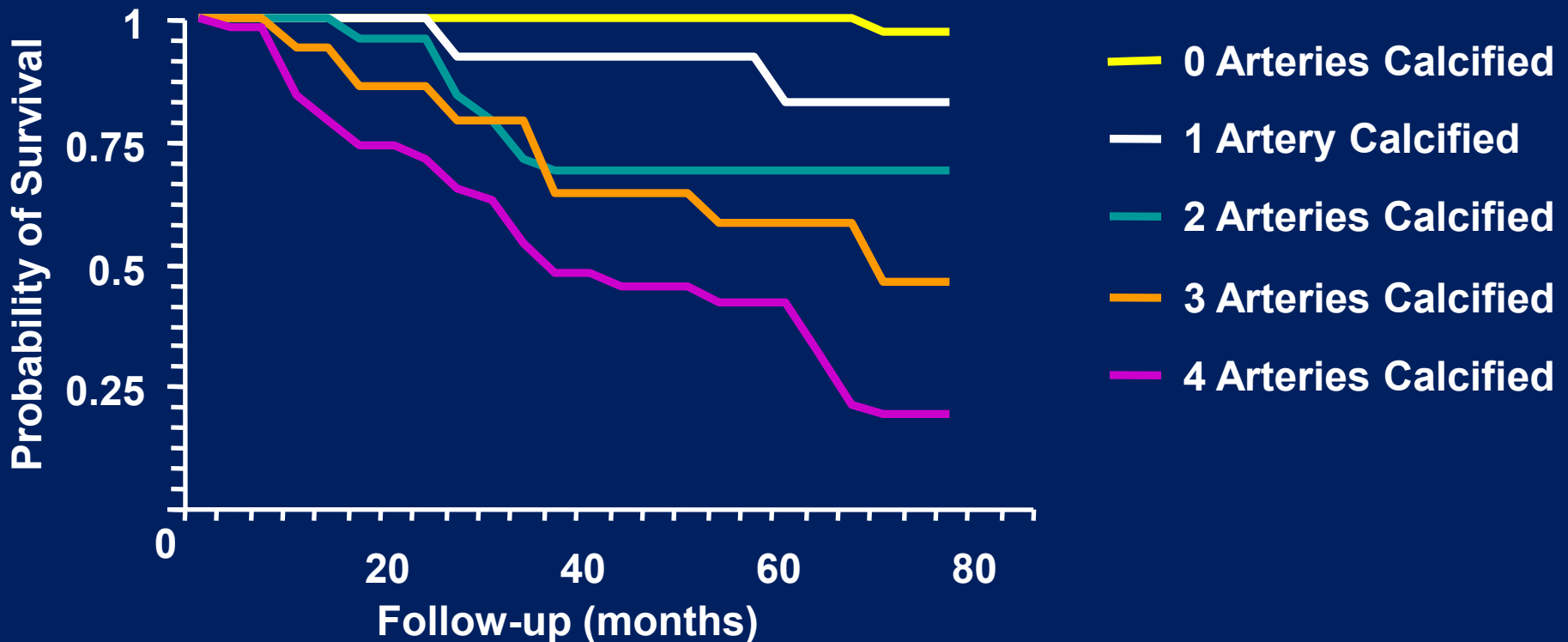


No. at risk

CCS = 0	46	42	42	39	34	18	4
CCS < 400	42	41	40	36	32	14	1
CCS ≥ 400	39	37	35	31	26	15	4

Block GA, et al. *Kidney Int.* 2007;71:438-441.

Probability of Survival Decreases With Increasing Arterial Calcification

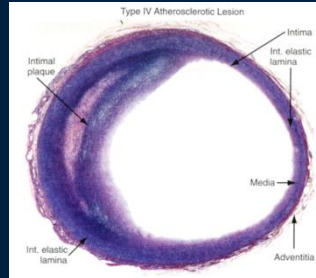


N=110 stable dialysis patients with ESRD.

$P < .0001$ comparison among groups.

Adapted from Blacher J et al. *Hypertension*. 2001;38:938-942.

Arterial Calcification: Intima vs Media



Intima

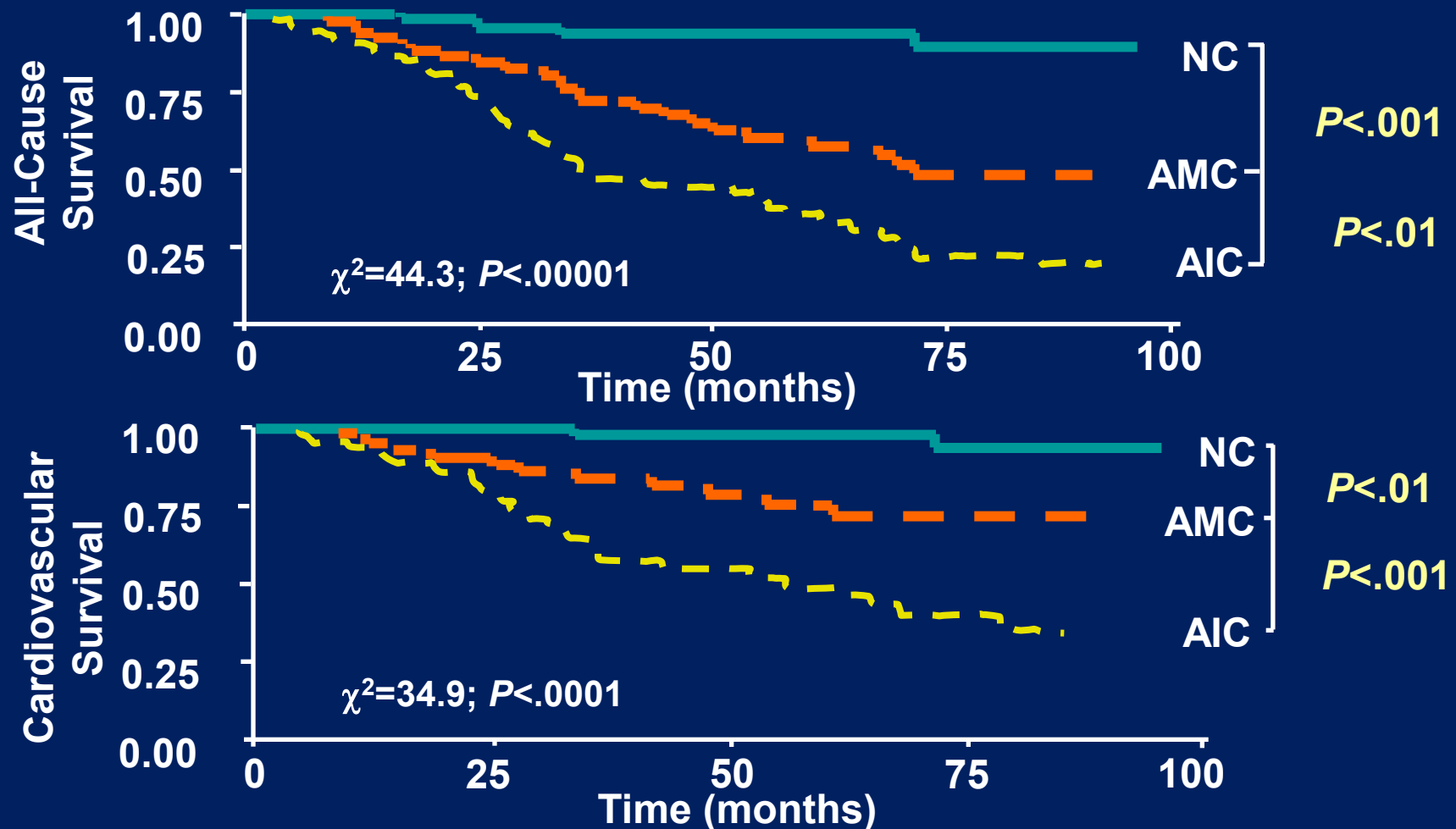


Media

Histology	Atherosclerosis Diffuse punctate morphology Aggregates of calcium crystals	Arteriosclerosis Linear deposits along elastic lamellae At most severe, a dense circumferential sheet of calcium deposition
Consequence	Acute closure (occlusion)	Vascular stiffness (nonocclusive)
Occurrence	Generalized CV disease	CKD, diabetes, aging (Monckeberg's sclerosis)
Factors	Lipid, macrophages, VSMC, inflammation	Elastin, VSMC

CKD Patients Often Exhibit Both Intimal and Medial Calcification

Impact of Arterial Calcification in Stable Hemodialysis Patients With CKD Stage 5



Bone Disease

Definition of Renal Osteodystrophy

- Alteration of bone morphology in patients with CKD characterized as abnormalities in bone turnover, mineralization, volume, linear growth, or strength.
- It is one measure of the skeletal component of the systemic disorder of CKD-MBD that is quantifiable by bone histomorphometry.

Role of PTH in Development and Progression of Renal Osteodystrophy

Low turnover

High turnover

Low

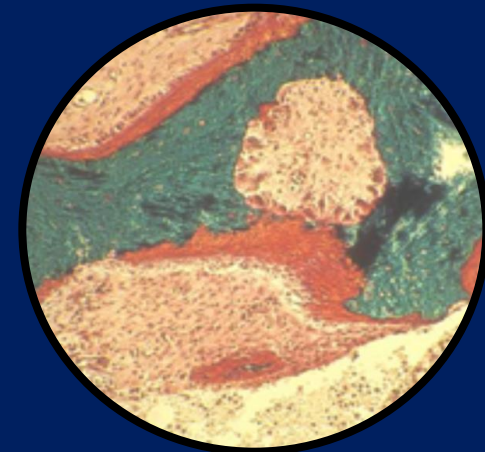
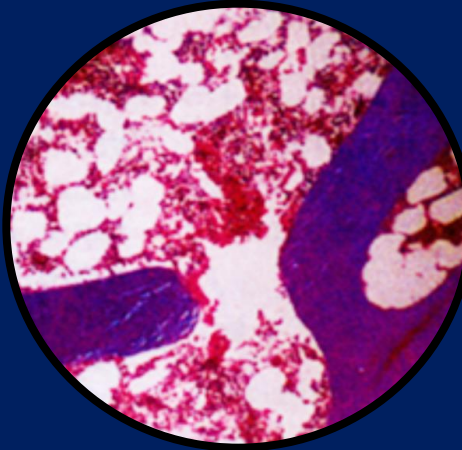
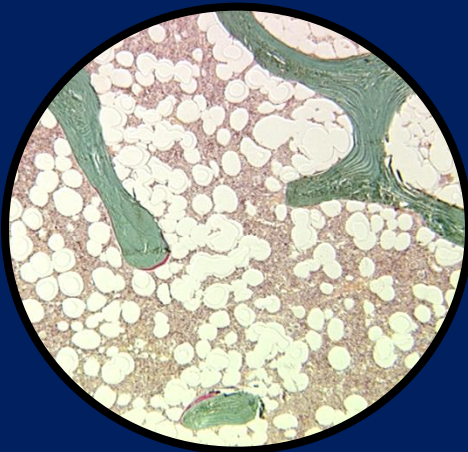
PTH

High

Adynamic
Osteomalacia

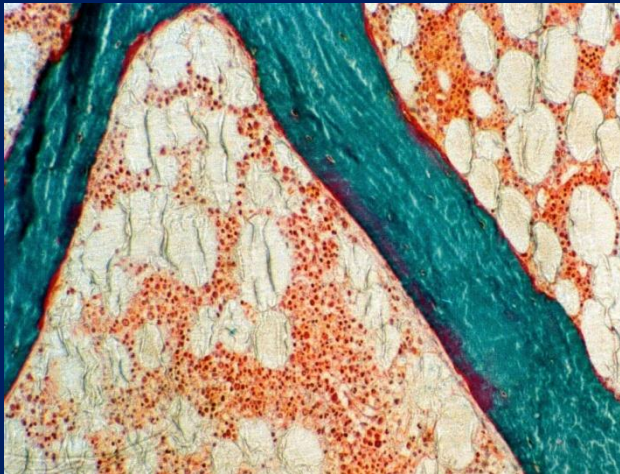
Normal Bone
Formation

Osteitis Fibrosa

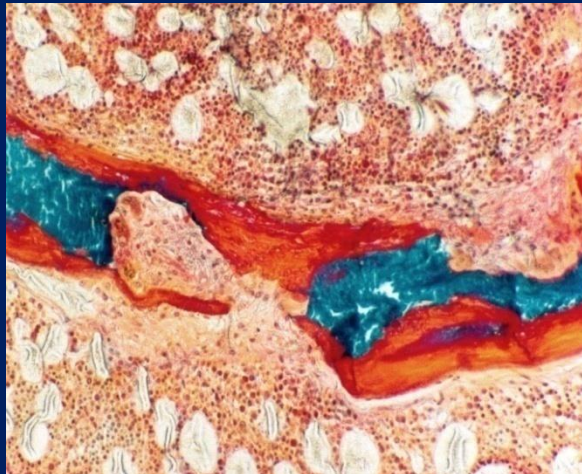


Mixed Lesion

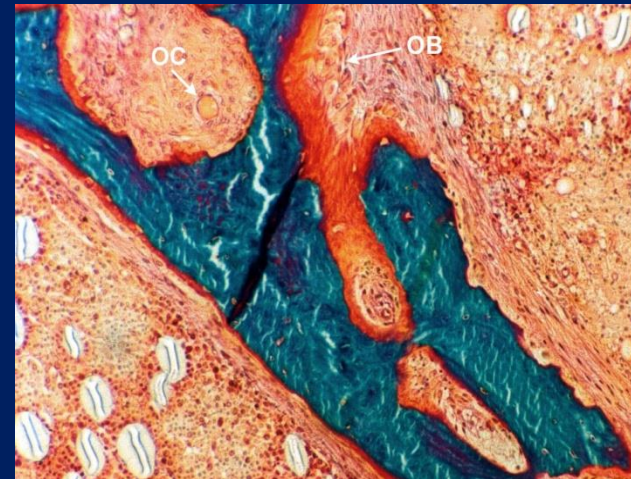
TMV Histologic Classification of ROD (KDIGO)



Adynamic bone



Mixed lesion



Osteitis fibrosa cystica

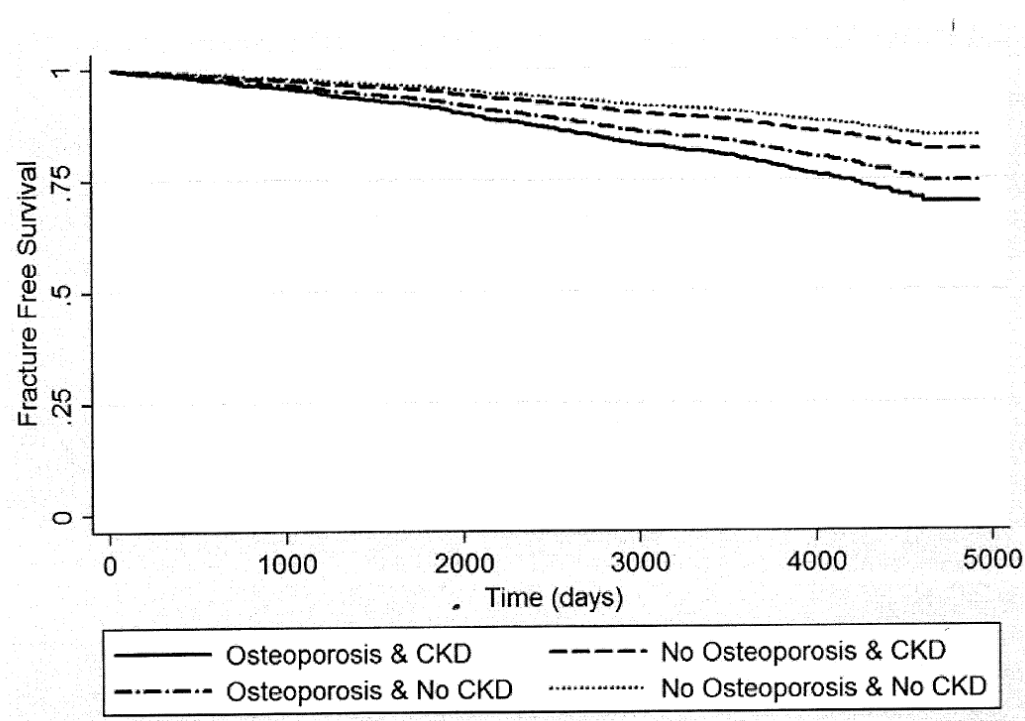
Turnover:	Low	High	High
Mineralization:	Abnormal	Abnormal	Abnormal
Volume:	Low	Low	Increased

TMV Distribution of Bone Lesions in CKD Stage 5

Mineralization (mlt_assm/Oth)	Turnover (bfrbs_assm)	Volume (bvtv_assm)	N (%)
Abnormal	1: Low	1: Low	166 (24.95)
		2: Normal	55 (11.25)
		3: High	46 (9.41)
	2: Normal	1: Low	35 (7.16)
		2: Normal	32 (6.54)
		3: High	16 (3.27)
	3: High	1: Low	25 (5.11)
		2: Normal	21 (4.29)
		3: High	10 (2.04)
Normal	1: Low	1: Low	8 (1.64)
		2: Normal	7 (1.43)
		3: High	5 (1.02)
	2: Normal	1: Low	17 (3.48)
		2: Normal	18 (3.68)
		3: High	2 (0.41)
	3: High	1: Low	4 (0.82)
		2: Normal	10 (2.04)
		3: High	74 (13.45)

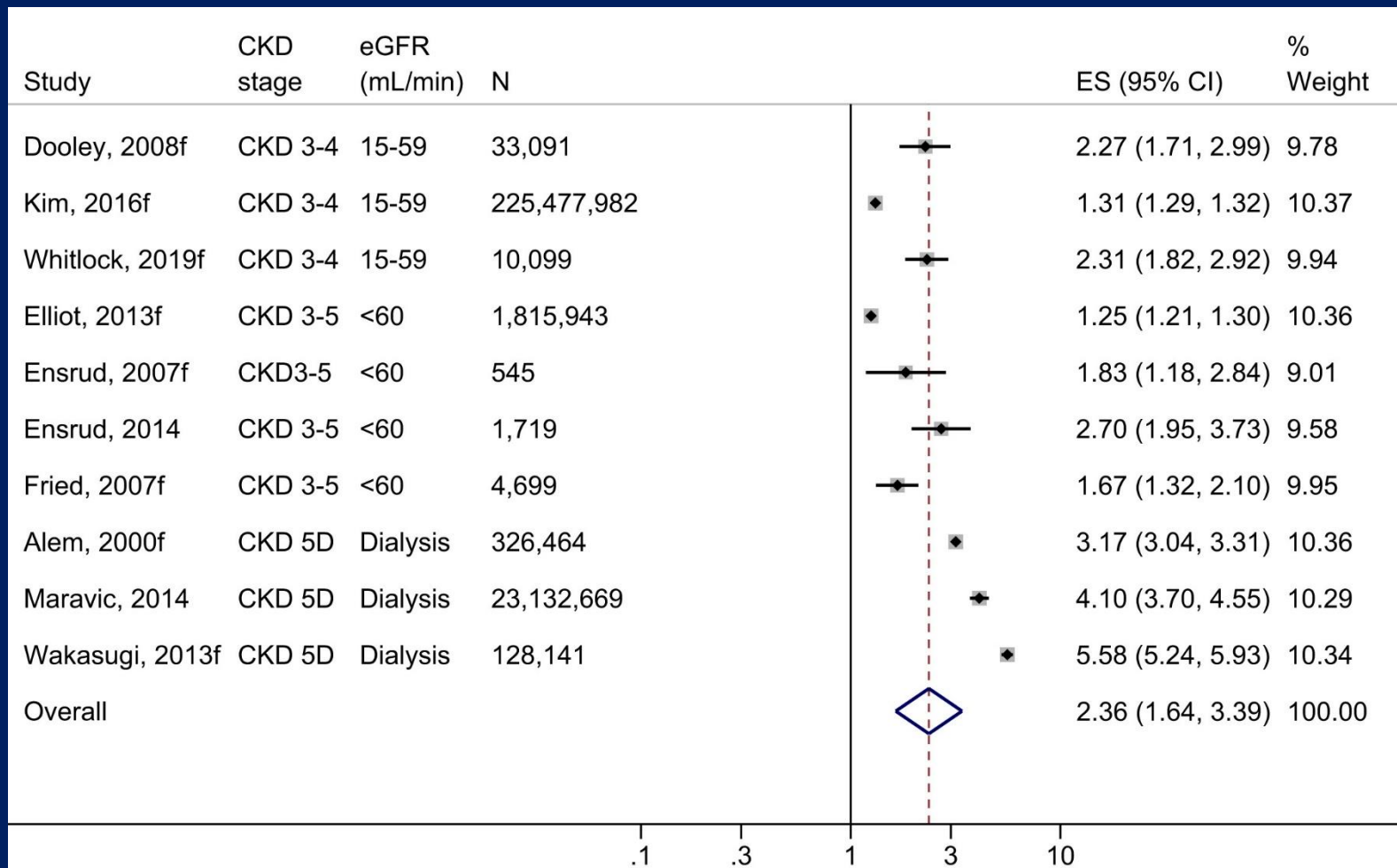
Bone Mineral Density and Fracture Risk with CKD - Corrected

Figure 3.

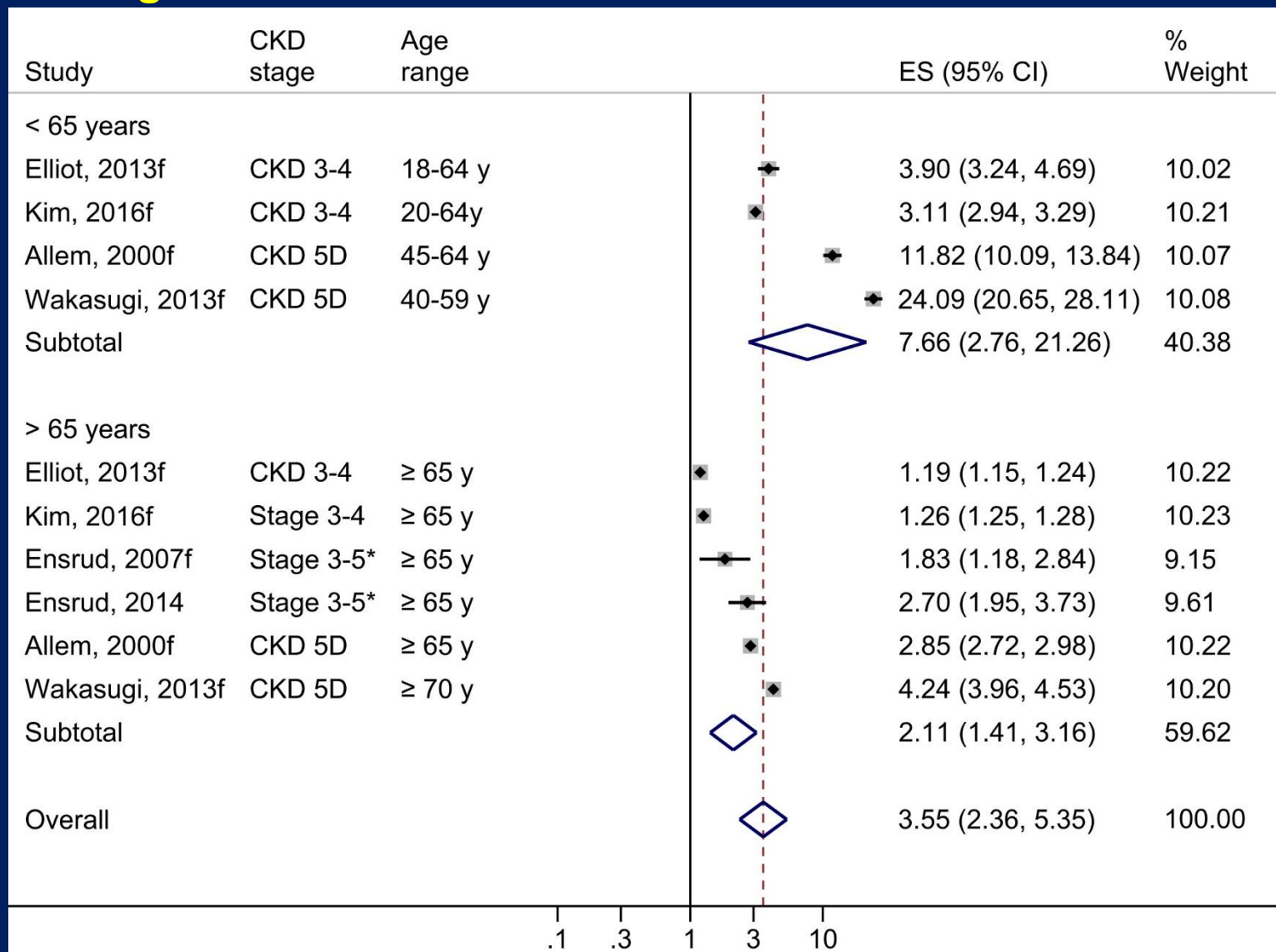


Fracture-free survival curves adjusted for age, race, sex, body mass index, hyperparathyroidism, and vitamin D deficiency. The relationship with fracture risk is demonstrated between the CKD and osteoporosis groups.

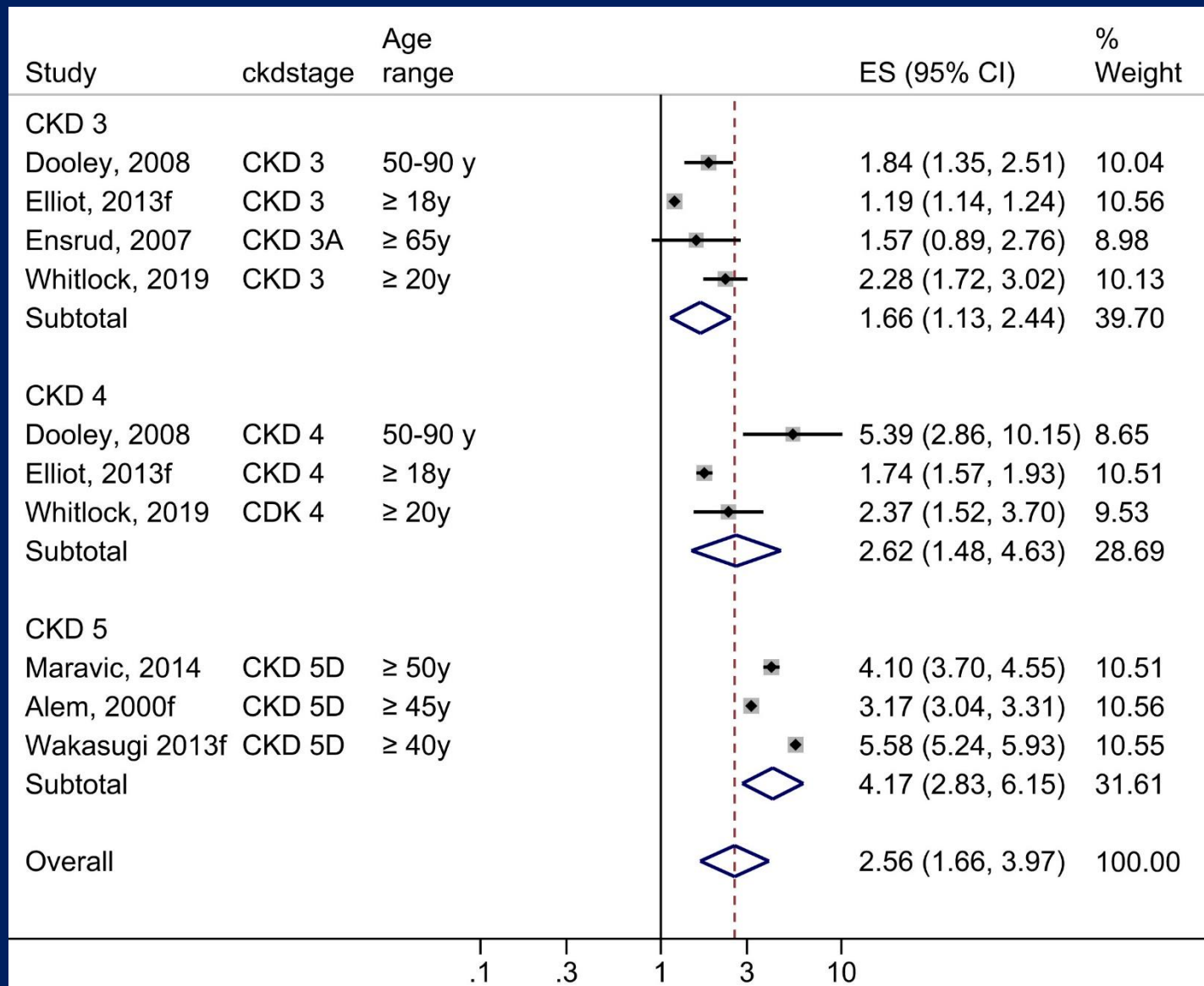
Overall Risk for Hip Fractures in CKD



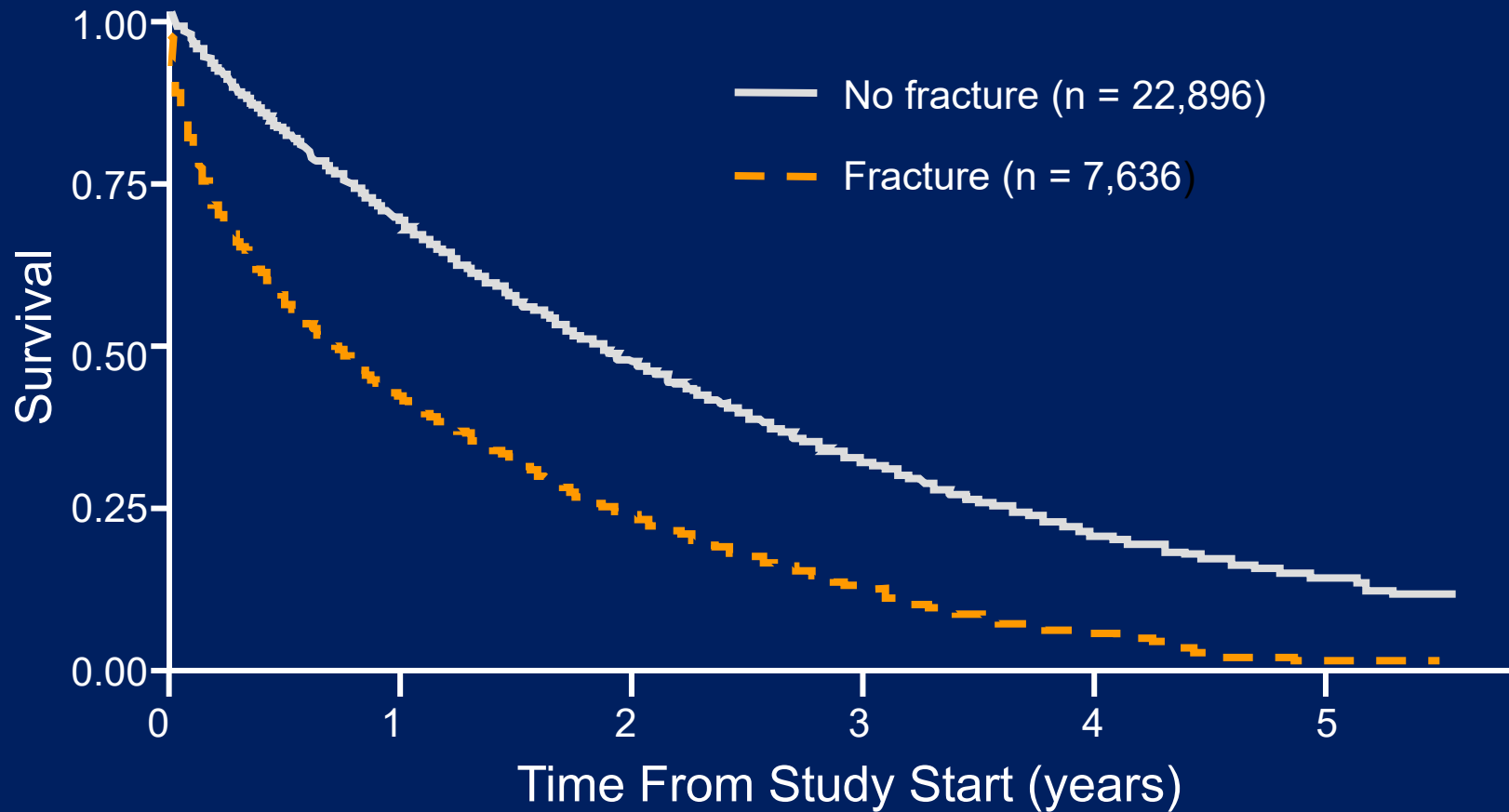
Hip Fracture Risk in CKD: Subgroup Analysis by Age Range



Hip Fracture Risk in CKD: Subgroup Analysis by Stage



Impact of Hip Fracture on All-Cause Mortality Risk in Dialysis Patients



CI = confidence interval.

Mittalhenkle A, et al. *Am J Kidney Dis.* 2004;44:672-679., Kaneko TM, et al. *Clin Orthop.* 2007;457:188-193.

Bone Turnover Markers

	High Turnover	Low Turnover
PTH	High	Low
Formation		
Total ALP	High	Low
BALP	High	Low
Intact PINP	High	Low
Resorption		
TRACP5b	High	Low

PTH – parathyroid hormone

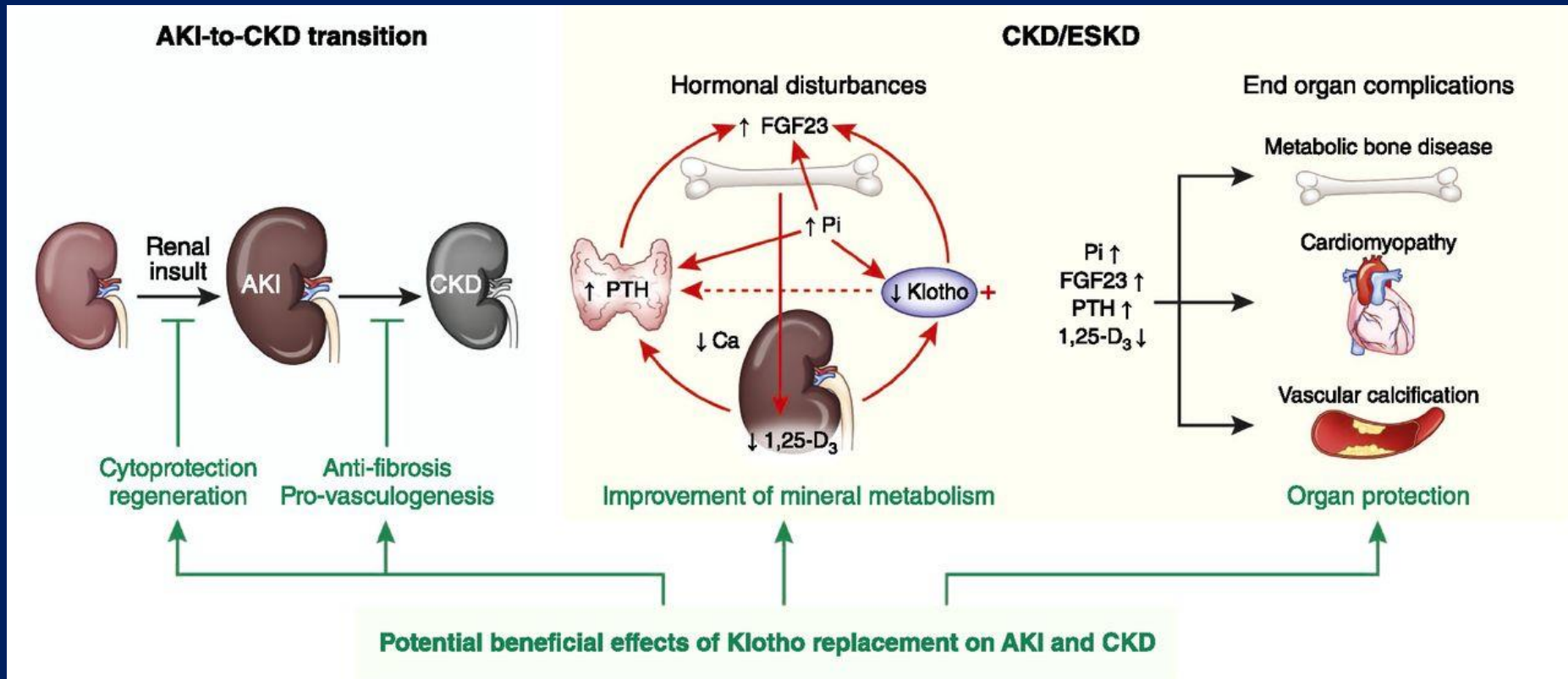
ALP – alkaline phosphatase

BALP – bone-specific alkaline phosphatase

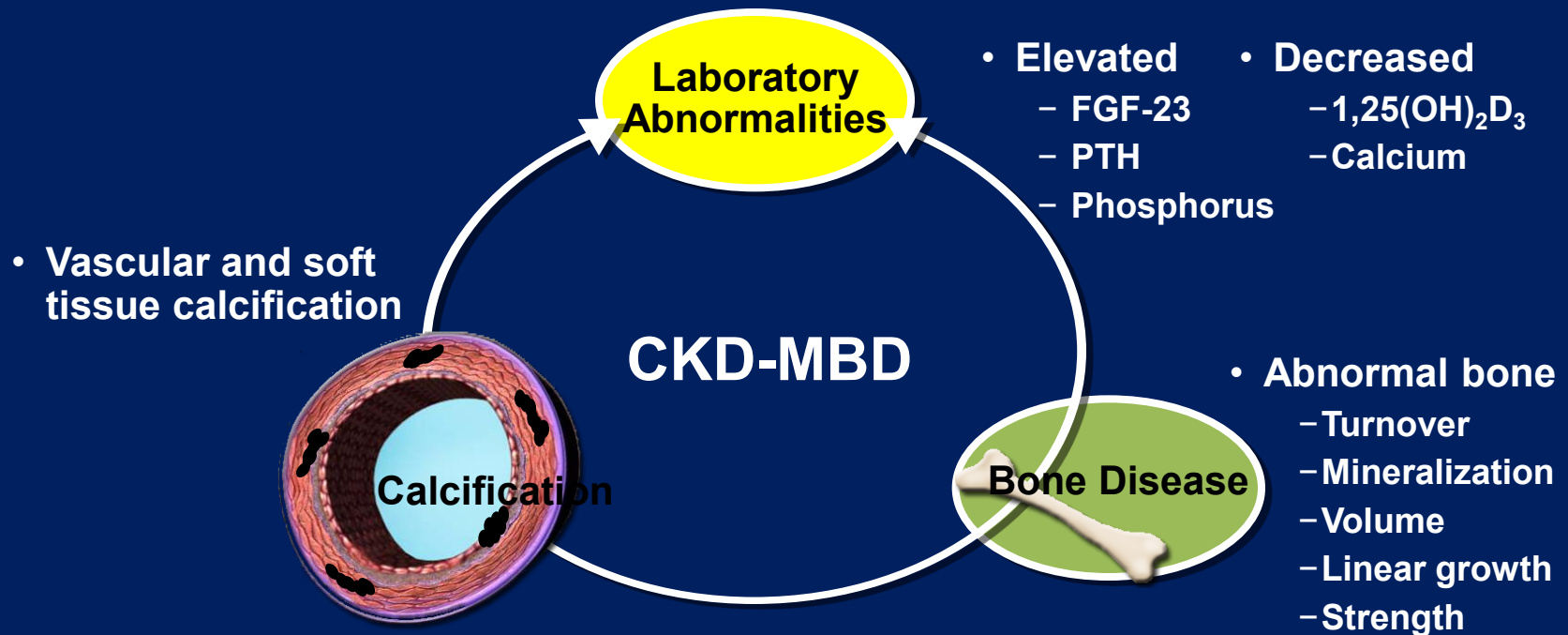
PINP – intact procollagen type 1 N-propeptide

TRACP5b – tartrate-resistant acid phosphatase isoform 5b

Role of Klotho in kidney and cardiovascular protection.



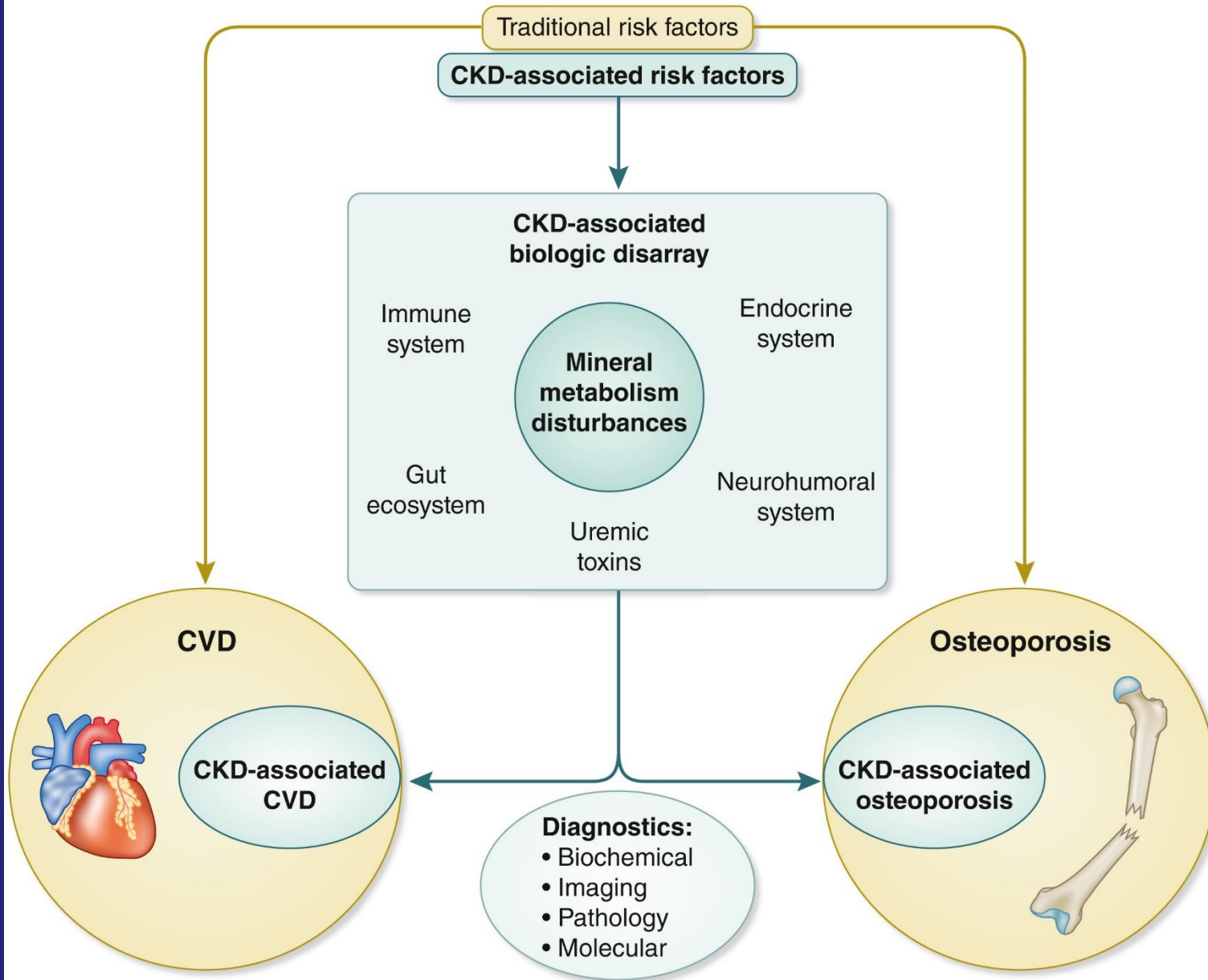
Chronic Kidney Disease-Mineral Bone Disorder



PTH = parathyroid hormone;
FGF-23 = fibroblast growth factor-23.

Kidney Disease: Improving Global Outcomes (KDIGO) CKD-MBD Work Group. *Kidney Int.* 2009;76

New conceptual framework moving towards personalized care in adults with CKD-MBD



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Questions ???



School of Athens / Scuola di Atene : Raphael / Raffaello Sanzio (1483-1520). Vatican. *The School of Athens* portrays Plato, Aristotle, and other ancient philosophers engaged in philosophic inquiry.